Paediatric safety in primary care: A cross-sectional mixed methods study of national incident report data

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Summary

Primary care is responsible for the majority of children's healthcare contact, yet there is a dearth of research into the safety of care provided to children in this setting. Confidential Enquiries highlight the need for improved vaccination, better recognition of seriously unwell children, and improved management of children with chronic conditions. This thesis therefore aimed to explore deficiencies in the vaccination process and in the primary care provided to 'unwell' children.

A cross-sectional mixed methods study of paediatric safety incidents involving vaccination or 'unwell' children, from primary care between 2002-2013 was conducted. The free-texts of 3913 reports submitted to the National Reporting and Learning System were classified to describe: incident types, contributory factors, incident outcomes, and severity of harm outcomes. Additionally, a literature review was conducted to identify potential interventions to address problem areas identified.

Key vaccination-related failures included vaccination with the wrong number of doses, at the wrong time, or with the wrong vaccine. Documentation failures and staff mistakes frequently underpinned these incidents, and vulnerable groups appeared more prone to incidents.

Key incidents involving 'unwell' children were related to: medication provision; and failures of diagnosis, assessment, referral, and communication, primarily related to telephone assessments. Medication errors were often the result of staff mistakes and failing to follow protocols. Incidents related to telephone assessment of 'unwell' children were often precipitated by protocol problems such as failing to assess children using the appropriate protocol.

The findings presented in this thesis provide an overview of paediatric safety problems in primary care, in addition to offering recommendations for improvement. Example recommendations include building IT infrastructure to address vaccination-related documentation discrepancies; electronic transmission of prescriptions to community pharmacies to reduce dispensing errors; and adapting clinical decision software to improve paediatric telephone-based assessments. The hypotheses generated from this work will form the basis of future work.

Abbreviations

BCG Bacillus Calmette-Guérin
BNF British National Formulary

CDS Computerised Decision Support

CPOE Computerised Physician Order Entry

DTaP/IPV Diphtheria Tetanus acellular Pertussis/Inactivated Polio

DTaP/IPV/Hib Diphtheria Tetanus acellular Pertussis/Inactivated Polio/ Haemophilus influenza type B

GP General Practitioners

Hib Haemophilus influenza b

Hib/Men C Haemophilus influenza b/Meningitis C

HIV Human Immunodeficiency Virus

HMIC Health Management Information Consortium

HPV Human Papilloma Virus

ICD-10 International Classification of Diseases - 10

ITS Interrupted Time Series

LINNAEUS Learning from International Networks About Errors and Understanding Safety

Men C Meningitis C

MeSH Medical Subject Headings

MMR Measles Mumps Rubella
NHS National Health Service

NICE National Institute for Health and Care Excellence

NPSA National Patient Safety Agency

NRLS National Reporting and Learning System

PCV Pneumococcal Conjugate
PISA PrImary care patient SAfety

PRN Pro Re Nata

RCPCH Royal College of Paediatrics and Child Health

RCT Randomised Controlled Trial

SIGN Scottish Intercollegiate Guidelines Network
Td/IPV Tetanus Diphtheria and inactivated Polio

UK United Kingdom

USA United States of America
WHO World Health Organisation

Glossary

	T
Adverse drug	Unintended, undesirable, or unexpected effects of prescribed
reaction	medications or of medication errors ¹
Adverse event	An undesired patient outcome that may or may not be the result of an
	error ²
Avoidable factors	Where there were identifiable failure in the child's direct care by an
	agency with direct responsibility for a child; where there were latent,
	organisational, or other indirect failures within one or more agency with
	direct or indirect responsibility for a child, where there was a failure of
	design, dilapidation or barriers, or inadequate maintenance by agencies
	with responsibility for public safety ³
Care failures	Failures in the healthcare provided directly to the child by an agency
	(including parents) with direct responsibility for that child ²
Caregiver	An individual who has responsibility for a child (such as parents)
Child health	A state of complete physical mental and social well-being not merely
	the absence of disease or infirmity, in children ¹
Chronic condition	A condition which typically requires follow up by health services,
	including repeated hospital admissions or outpatient appointments,
	long-term medication use, or use of support services such as
	physiotherapy ⁴
Contributory factor	A circumstance, action, or influence that is thought to have played a
	part in the origin or development of an incident or to increase the risk
	of an incident ¹
Contributory incident	A patient safety incident that played a part in the origin or development
	of another incident or increased the risk of another incident
Error	Deviation in a process of care that may or may not cause harm to
	patients ¹
Error of commission	An error that occurs as a result of an action taken ¹
Error of omission	An error that occurs as a result of an action not taken ¹
	1

Joint Commission Resources 2005
 World Health Organisation 2009
 Pearson G 2008
 Hardelid P, et al. 2013

Epistemology	The study of knowledge, what constitutes knowledge, and how we
	obtain it ⁵
EU 15+	The 15 countries of the European Union prior to 2004 plus Australia,
	Canada and Norway ⁶
Excess deaths	The difference between the number of deaths observed and the number
	of deaths that would have occurred given the same death rate as a
	comparable country
First-access services	The services to which 'unwell' patient initially present (includes primary
	and emergency care)
Global Trigger Tool	A method to measure all-cause harm using patient records ⁷
Harm	A negative patient outcome that includes temporary or permanent
	impairment of the physical, emotional, or psychological function or
	structure of the body and / or pain requiring intervention ¹
Hazard	A situation or event that introduces or increases the probability of an
	adverse event arising from a danger or peril, or that increases the
	extent of an adverse event ¹
Healthcare harm	Harm arising from, or associated with, plans or actions taken during the
	provision of healthcare rather than an underlying disease or injury ¹
Human error	One category of potential causes for unsatisfactory activities or
	outcomes ¹
Human factor	Study of the interrelationships between humans, the tools, equipment,
	and methods they use, and the environments in which they live and
	work ¹
latrogenic	An illness or injury resulting from a diagnostic procedure, therapy, or
	other element of healthcare ¹
Incident type	A descriptive term for a category made up of incidents of a common
	nature grouped because of shared, agreed features ¹
Incident outcome	The impact upon a patient or organisation which is wholly or partially
	attributable to an incident ¹
Looked-after child	A child under the care of the local authority
Modifiable factors	Extrinsic factors which could potentially be addressed, that are not
	necessarily related to healthcare ²
Near miss	Events or situations that could have resulted in an accident, injury, or

Bourgeault I, Dingwall R and DeVries R 2010
 Viner RM et al. 2014
 Parry G, Cline A and Goldmann D 2012

	illness, but did not, either by chance or through timely intervention ¹
Ontology	The study of reality and how we understand existence ⁴
Patient safety	An event or circumstance which could have resulted, or did result, in
incident	unnecessary harm to a patient ¹
Preventable harm	A negative patient outcome that would not have occurred if the patient
	had received ordinary standards of care appropriate for the time ¹
Primary care	The provision of integrated, accessible health care services by clinicians
	who are accountable for addressing a large majority of personal health
	care needs, developing a sustained partnership with patients, practicing
	in the context of family and community ⁸
Pro re nata (PRN)	Take medication as required
Quality of care	The degree to which health services are timely, efficient, equitable,
	safe, patient-centred, and effective ⁹
Recursive	Repeated application of a rule, definition, or procedure to results
Root cause analysis	A systematic process of investigating incidents to identify the multiple,
	underlying, and latent contributory factors ¹
Substandard care	Failure to apply the principles and practices accepted by a healthcare
	profession, as expected ¹
Systems error	An error that is not the result of an individual's actions, but the
	predictable outcome of a series of actions and factors that comprise a
	diagnostic or treatment process ¹
Unsafe care	Failure to make evidence-based clinical decisions to maximise the
	healthcare outcomes of an individual and failure to minimise the
	potential for harm ¹
Unwell child	A child with signs, symptoms, diagnoses, or prescribed medications
	implying illness

 ⁸ Institute of Medicine 1996
 9 Institute of Medicine 2001

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Chapter 1: Introduction

The United Kingdom (UK) has one of the worst child mortality rates in Western Europe and the contribution of unsafe healthcare to this high mortality is unknown (Carson-Stevens A et al. 2015; Viner RM et al. 2014). However 26% of child deaths have identifiable failures in care, and primary care is responsible for 90% of healthcare contact (Carson-Stevens A et al. 2015; Pearson G 2008). Issues with primary care quality are apparent through high rates of inappropriate hospital admissions and paediatric referrals, in addition to high mortality rates for diseases dependant on first-access services such as meningitis and pneumonia compared to other Western European countries (Wolfe I et al. 2011; Wolfe I et al. 2013).

This chapter will discuss the landscape of child health and quality improvement globally with particular focus on UK issues. Child health outcomes and associated care quality data will be presented. European care models will be discussed, compared, and contrasted with UK models. The importance of healthcare safety to child health will be reviewed, in addition to the various available methods for assessing the nature and burden of unsafe care.

1.1 Child health background

1.1.1 Epidemiology

Since the turn of the millennium substantial gains in child health have been achieved globally and there has been an accelerated decrease in child mortality. (Bryce J et al. 2013; Wang H et al. 2014). There has been a dramatic shift in the global burden of disease and the prevalence of communicable diseases has decreased (Bryce J et al. 2003; Bryce J et al. 2013; Liu L et al. 2012). The Millennium Development Goals, particularly goal 4 to decrease under-5 child mortality by 67% from 1990 to 2015, instigated this shift in disease burden. Although the Millennium Development Goals have not been fulfilled, they successfully accelerated global improvements in child health (Bryce J et al. 2013).

Child survival in Europe has mirrored improvement in less developed countries: consequently the landscape of child health in Europe has changed. Healthcare amenable deaths from infections are less of a problem whilst deaths from non-communicable diseases are an increasing burden on child health (Liu L et al. 2012; Wang H et al. 2014; Wang H et al. 2012). Non-communicable diseases are responsible for significant child morbidity and mortality, and they account for 79% of disability-adjusted life years lost (Liu L et al. 2012; Wang H et al. 2014; Wang H et al. 2012; Wolfe I et al. 2013). The Global Burden of Disease Study reports that respiratory diseases, neuropsychiatric disorders, congenital abnormalities, and musculoskeletal disorders have the greatest impact on childhood morbidity in Western Europe (Institute for Health Metrics and Evaluation 2010).

The UK has achieved considerable reductions in child mortality but these gains in child health have not matched those of comparable European countries i.e. the EU 15+ that are comparable in terms of health expenditure (Viner RM et al. 2014). This slower decline in child mortality has seen the UK drop from having one of the lowest rates of child mortality rates in Europe to having the highest rate in Western Europe (Viner RM et al. 2014). Annually around 6000 infants, children, and adolescents die in England and Wales, which equates to almost 2000 excess deaths compared to Sweden. Reductions in mortality have largely been in children aged 1-12 years, consequently most deaths (67%) occur in infancy, followed by adolescence (Sidebotham P et al. 2014a; Sidebotham P et al. 2014b; Wolfe I et al. 2014).

1.1.2 Child health outcomes and care quality

The UK performs sub-optimally in numerous measures of child health and wellbeing. Over a quarter of child deaths reviewed in a 2006 Confidential Enquiry had identifiable failures in care, 43% had potentially avoidable factors, and 21% of deaths had modifiable factors considered healthcare amenable i.e. potentially preventable (Pearson G 2008; Wolfe I et al. 2014).

Up to 30% of deaths in infants and adolescents, when mortality peaks, are thought to have modifiable factors (Wolfe I et al. 2014). The high UK infant mortality rate, relative to comparable European countries, likely reflects high rates of preterm birth that are considered preventable and a reflection of the quality of midwifery, obstetric and newborn care (Wolfe I et al. 2014). Viner RM et al. 2014 estimate that such failures culminate in approximately 1000 excess annual infants deaths compared to the European average.

Chronic conditions (e.g. mental health conditions, cancer, respiratory conditions) are estimated - by a recent epidemiological review of UK child deaths using routinely collected vital statistics and administrative health care data - to be responsible for 60-70% of child deaths (Hardelid P et al. 2013). Deaths from chronic conditions pose a considerable problem for the UK and have been identified as a priority area for improvement by child health reviews (Hardelid P et al. 2013; Wolfe I et al. 2014). Mortality from endocrine, respiratory, digestive, and neuropsychiatric disorders is higher in the UK than comparable European countries, and deaths from such non-communicable diseases are not decreasing in line with comparable countries (Institute for Health Metrics and Evaluation 2010; Viner RM et al. 2014; Wolfe I et al. 2014) There is also increasing epidemiological evidence from analyses of World Health Organisation (WHO) mortality data that cancer survival is worse and diagnosed later in the UK (Viner RM et al. 2014; Wolfe I et al. 2014).

A case note review to determine the quality of care delivered to children with diabetes identified that fewer than 6% of children in England receive evidence based care in line with published guidance, thus resulting in poor diabetic control, preventable emergency admissions, and preventable deaths (National Diabetes Audit 2008). Consequently children in England and Wales have poorer diabetic control than children in comparable European

countries: 83% have HbA1c concentrations above target levels and 9% experience diabetic ketoacidosis annually (National Diabetes Audit 2008; Wolfe I et al. 2011).

National UK audits of care quality for childhood epilepsies, using 12 quality indicators, highlighted similar failures: 35% of children did not have an appropriate initial assessment, and access to specialist paediatric neurologists and nurses was not universal (Royal College of Paediatrics and Child Health 2012).

Considerable geographical variation health outcomes exist in the UK and these may reflect variation in care quality. Emergency admissions for children with asthma varied from 25.9/100 000 population in some locations to 641/100,000 in others (Asthma UK 2007). The UK has a higher asthma-related mortality than comparable countries (World Health Organisation Regional Office for Europe 2012). This has been a long-standing problem, and in 2000 the UK was in the lowest EU 15+ quartile for child mortality from non-communicable diseases across all ages (Viner RM et al. 2014; Wolfe I et al. 2011).

Poor primary care quality impacts negatively on secondary and emergency care services. In the UK 75% of asthma-related admissions are deemed avoidable with better primary care. A third of short stay admissions in infants occur for minor illness, which should ideally be managed in the community, and 36% of referrals to paediatricians are deemed inappropriate (Asthma UK 2007; Milne C et al. 2010; Saxena S et al. 2009).

Death rates from diseases that rely heavily on first-access services are arguably a reflection of primary care quality. Compared to other European countries, the UK has high rates of mortality from illnesses that rely heavily on first-access services such as meningococcal infections and pneumonia (Wolfe I et al. 2011; Wolfe I et al. 2014; World Health Organisation Regional Office for Europe 2012). Unsafe care may be partially responsible these results suggesting poor care quality in the UK, and this will be discussed further in section 1.1.3.

1.1.3 Inequalities in child health

Considerable inequalities in health persist in high-income countries particularly in children (Mackenbach JP et al. 2008; Sidebotham P et al. 2014a). This is demonstrated by the variation in child survival by country and population demographic. Differences in child health outcomes are vast within and between countries in Western Europe. Suggested explanations for those variations include: the organisation and quality of healthcare services, social inequalities, and cultural and economic factors (Sidebotham P et al. 2014a).

Key social determinants of health include poverty, inequality, and social policies (Marmot M et al. 2012; World Health Organisation Commission on Social Determinants of Health 2008). Child mortality is not only associated with absolute poverty, but also relative poverty and inequality i.e. imbalanced distribution of wealth (Adamson P 2012; Collison D et al. 2007; Marmot M et al. 2012; Pritchard C and Williams R 2011; Wolfe I et al. 2014; Wolfe I et al. 2013). For example, the Nordic countries with the lowest proportion of impoverished households in Europe have the lowest child mortality rates; and the five high-ncome countries with the highest child mortality rates also have the highest inequalities in household income (Collison D et al. 2007; Pritchard C and Williams R 2011; Sidebotham P et al. 2014a; Wolfe I et al. 2014).

The inverse association of socioeconomic status and childhood mortality is well acknowledged and relative poverty has been demonstrated as a key determinant of child deaths in the United States of America (USA), Australia, and New Zealand, as well as in the UK (Arntzen A and Nybo-Andersen AM 2004;Blakely T et al. 2003;Freemantle N et al. 2009;Gakidou E et al. 2010;Petrou S et al. 2014;Pickett KE and Wilkinson RG 2007;Rodwin VG and Neuberg LG 2005;Sidebotham P et al. 2014a;Spencer N 2004). Poor socioeconomic status has been associated with child mortality from a range of causes such as suicides, poisoning, and cancer, in particular acute lymphoblastic leukaemia as well as sudden infant death syndrome (Agerbo E et al. 2002;Edwards P et al. 2006;Kong KA et al. 2010;Lightfoot TJ et al. 2012;Pickett KE and Wilkinson RG 2007;Sidebotham P et al. 2014a;Spencer N 2004;Wood AM et al. 2012).

When child poverty is defined as the number of households, with children, in which the disposable household income is less than 50% of the median disposable household income for that country, the UK has one of the highest rates of child poverty among high income

countries (Adamson P 2012). However, it is worth noting that numerous definitions of child poverty exist and that the UK has high rates of relative poverty rather than absolute poverty (Adamson P 2012). Currently 35% of UK households with children are estimated to have insufficient income for acceptable standards of living (Padley M and Hirsch D 2014; Wolfe I et al. 2015). Child mortality statistics reflect the high rates of relative poverty and the marked social inequalities present in the UK (Wolfe I et al. 2014) Standards of living are predicted to worsen, due to reduced public service funding and continued economic pressures (Wolfe I et al. 2015). Therefore, the importance of access to high quality healthcare is argued to become ever more apparent to mitigate further deterioration of child health outcomes (Viner RM et al. 2014; Wolfe I et al. 2014).

1.1.3.1 Inverse care law

Over 40 years ago Tudor Hart described the inverse care law: provision of "good medical care tends to vary inversely with the need for it in the population served" (Hart JT 1971). This phenomenon has persisted and children tend to suffer disproportionately, as childhood is a particularly vulnerable period (Webb E 1998). The inverse care law is clearly visible in children of vulnerable and marginalised populations such as: refugees, travellers, and ethnic minority groups that have above average mortality (Alio AP et al. 2010;Anachebe NF 2006;Sidebotham P et al. 2014a;Webb E 1998;Wolfe I et al. 2011). For example rates of mortality, preterm birth, and communicable diseases in Roma children far exceed those of the general population across Europe (Wolfe I et al. 2013). A range of complex and interacting factors have been proposed to underlie these trends including access to healthcare, socioeconomic disadvantage, poor living conditions, higher rates of consanguinity, and genetic predispositions (Alio AP et al. 2010;Sidebotham P et al. 2014a;Smith GD 2000).

It has been noted that children in refuges who greatly need healthcare have poor access to it (Webb E et al. 2001). Their child health records tend to be incomplete, and developmental screening and vaccination uptake in this population is sub-optimal (Webb E et al. 2001). In addition they have unmet mental health needs compounded by their past experiences, which are often violent and abusive (Webb E et al. 2001).

1.1.4 Child health services

Since the 20th century the relative contribution of healthcare to population health has increased as living conditions improved in response to industrialisation (Wolfe I et al. 2014). Access to high quality healthcare can modify the negative effect of biological, social, cultural, and financial factors on child health (Sidebotham P et al. 2014a; Wolfe I et al. 2014).

A unique and equitable attribute of the UK National Health Service (NHS) is that it is free at the point of delivery, and 90% of contact with healthcare providers occurs via primary care services delivered in community settings. Countries with successful primary care systems tend to achieve better population health outcomes (Institute of Medicine 1996;Starfield B 1991;Starfield B 1994;Starfield B et al. 2005). There is no consensus on the optimum organisation of primary care for children and numerous models exist.

Primary care for children differs considerably between European countries in terms of organisation, the provision of out-of-hours care, the professionals providing care, and the training that care providers receive (Wolfe I et al. 2013). The organisation and therefore the role of primary care can range from acting as a gateway to specialist services and paediatricians, as in the UK and the Netherlands, or access to paediatricians may be unlimited as in Sweden (Ahgren B 2003;Schäfer W et al. 2009;Wolfe I et al. 2011;Wolfe I et al. 2013). Out-of-hours care has become increasingly centralised in many European countries with groups of general practitioners (GPs) providing out-of-hours care and telephone triaging becoming more prevalent. Differences in the organisation of first-access services, including out-of-hours care, likely accounts for the large variation in rates of inappropriate emergency admissions (Ahgren B 2003;Wolfe I et al. 2011;Wolfe I et al. 2013).

In the UK first-access services, within primary care, are primarily (although not exclusively) the responsibility of and delivered by GPs, whereas in Germany and France they are delivered predominantly by primary-care-based paediatricians, and in Sweden they are delivered by co-located and collaborating GPs and primary-care-based paediatricians (Ahgren B 2003;Nolte E and McKee M 2008;Wolfe I et al. 2011;Wolfe I et al. 2013). This collaboration includes nurse-led care, which has become the norm for the management of certain conditions like asthma in Sweden. There are considerable differences in healthcare professionals' training between countries: for example, GPs in

Sweden undergo at least three months specialist training in paediatrics whereas in the UK GPs may undergo no postgraduate paediatric training (Ahgren B 2003; Wolfe I et al. 2011; Wolfe I et al. 2013).

The organisation of health services for children largely influences how chronic conditions are managed, for example in the community or hospital setting, and by paediatricians, nurses, or GPs (Wolfe I et al. 2011; Wolfe I et al. 2013). There is a relative paucity of policy directives in European countries on the management of chronic paediatric conditions compared with adult care (Wolfe I et al. 2013). Consequently large variations exist between countries in the delivery of healthcare to children with chronic conditions.

The Swedish model of paediatric primary care delivery has been argued to be the gold standard design; this comprises multidisciplinary teams including GPs, paediatricians, and children's nurses, co-located in primary care centres (Ahgren B 2003; Wolfe I et al. 2013). This model was designed to improve continuity of care and paediatric care quality by increasing multidisciplinary collaboration (Ahgren B 2003). In contrast the UK model is arguably the poorest for child health.

The quality of primary care for children has been neglected in the UK despite children accounting for over 20% of GP consultations (Gill PJ et al. 2011; Hippisley-Cox J et al. 2007; Royal College of General Practitioners 2008). The pay for performance quality indices which incentivise disease management in general practice almost completely overlook management of paediatric diseases: <3% of quality and outcome framework (QOF) indicators for general practices in England and Wales are relevant to children (Wolfe I et al. 2013). Wolfe I et al. 2013 also criticise the system for incentivising competition and professional self-interest rather than collaboration.

Despite a lack of consensus on how best to organise primary care to meet the needs of children, the epidemiological changes in the burden of disease and the child health needs warrant a change from the traditional hospital-centric model, designed to treat acute illnesses, to increasing care in the community and better equipping primary care to manage chronic conditions (Wolfe I et al. 2011;Wolfe I et al. 2013). The Healthcare Commission reports that 46% of UK trusts provide poor paediatric care in the community and that less focus is needed on hospital care and more on the management of chronic conditions in the community (Healthcare Commission 2007). The changing needs of the paediatric population must also be reflected in how primary care is funded, for example

Sweden, Italy, France, Germany, and the Netherlands have substantially more doctors per child than the UK (Royal College of Paediatrics and Child Health 2008; Wolfe I et al. 2011; World Health Organisation).

There is no gold-standard method of measuring the quality of child healthcare and the literature evaluating care quality to date in the UK has largely focused on disease-specific indicators, child death reviews, and mortality statistics. Use of mortality data as a surrogate measure of care quality has considerable limitations: the data are prone to numerous biases, mortality data lacks context, meaningful analysis of mortality data is difficulty - highlighted by the internal heterogeneity of the 'cause of death' categories used by Viner RM et al. 2014; and the data are likely confounded at least partially by unknown factors (Hardelid P et al. 2013;Hardelid P and Gilbert R 2013;Johnston BD 2014). Therefore, any conclusions about care quality originating from mortality data, or from comparing mortality data between countries, must be interpreted cautiously. The methods available to measure healthcare quality will be discussed further in Section 1.3.

1.2 Healthcare safety and children

1.2.1 Overview of the history of healthcare safety

High quality healthcare is defined by the Institute of Medicine as: safe, equitable, patient-centred, timely, efficient, and effective (Committee on Quality of Health Care in America Institute of Medicine 2001). The importance of high quality healthcare has been highlighted in section 1.1 and healthcare safety is an integral component of this. This section will discuss the background of paediatric safety including research conducted to date and current gaps in the evidence.

The principle of patient safety and healthcare harm is not new. Thomas Inman first coined the phrase "primum non nocere" also known as "first do no harm" in 1860; today this is considered a fundamental concept in medical ethics (Sokol DK 2013). Patient safety as a field however has only gained traction since the turn of the millennium, in response to high-profile healthcare scandals such as the Bristol Heart Inquiry in the UK, and the USA Institute of Medicine report 'To err is human'. (Kennedy I 2001; Kohn LT et al. 1999) The mortality rates of infants undergoing heart surgery in Bristol was double that of comparable hospitals in England (Kennedy I 2001). Similarly in the USA, annual mortality from healthcare errors was greater than from road traffic accidents, breast cancer, or Acquired Immune Deficiency Syndrome (Kohn LT et al. 1999).

Unfortunately two decades on, the landscape of patient safety has not changed significantly and healthcare tragedies continue, as evidenced by the Francis report on Mid-Staffordshire (Francis R 2013;Landrigan CP et al. 2010). The failures and recommendations for improvement published in the Francis report echo those of the Bristol Heart Inquiry. For example, healthcare professionals had been reporting and raising concerns about the safety of care provided in both the Bristol Royal Free in the early 1990s and more recently in Mid Staffordshire, long before the organisations themselves recognised and investigated these failings (Francis R 2013;Kennedy I 2001). Similarly in both cases data demonstrating above average mortality in the respective institutions were readily available, but the gravity of the situation was not acknowledged. It is therefore unsurprising that healthcare organisations are repeatedly criticised for their failure to learn (Francis R 2013;Kennedy I 2001).

Efforts to measure and improve patient safety have largely focused on the hospital rather than the community setting (Carson-Stevens A et al. 2015; Rees P et al. 2015). There has

also been little focus on paediatric safety, which is reflected in the absence of this topic from UK policy directives (Carson-Stevens A et al. 2015; Rees P et al. 2015).

1.2.2 Patient safety in child health

Children are particularly susceptible to unsafe care. Children who are very young, socially deprived, or have complex medical conditions are particularly vulnerable to adverse events (Sidebotham P et al. 2014a). They typically depend on parents or caregivers to recognise illness, take them to a healthcare professional, provide medical histories to physicians, and administer treatment (Wolfe I et al. 2011). Children also depend more on the people surrounding them to recognise unsafe care and to question the care they receive (Carson-Stevens A et al. 2015).

The epidemiology of disease, healthcare needs, and physiology of children differ significantly to those of adults. Numerous diseases are specific to childhood, and the effect of disease in children varies in terms of the signs and symptoms they present with and the speed at which they deteriorate (Walsh KE et al. 2014; Wolfe I et al. 2011). Differences in physiological reserve partially account for these differences (Walsh KE et al. 2014).

The epidemiology of disease, healthcare needs, and physiology, also vary by age, within the paediatric population (Wolfe I et al. 2011). The heterogeneity of this population-ranging from neonates to adolescents- poses safety challenges that are unique to the specialty of paediatrics. For example, children often require weight-based medication that can involve complex calculations requiring skill and familiarity with prescribing and administering medications, predisposing them to medication errors (Carson-Stevens A et al. 2015; Walsh KE et al. 2014).

The culminations of these vulnerabilities to unsafe care result in substantial estimates of iatrogenic harm in this population. Approximately 15-35% of hospitalised children are estimated to suffer an adverse event, and in the UK 26% of deceased children have identifiable failures in care (Pearson G 2008; Walsh KE et al. 2014). Estimates of the incidence of adverse events in children in primary care in the UK- where 90% of patient contact occurs-do not exist (Carson-Stevens A et al. 2015).

Medication errors have been estimated to be three times as common in children compared with adults (Department of Health 2000; Wong IC et al. 2009). Consequently these have been the main focus of the limited research into paediatric safety, although most studies focus on secondary care errors (Carson-Stevens A et al. 2015). Children appear particularly prone to tenfold medication errors, especially neonates and those with chronic conditions requiring complex treatment (Doherty C and Mc Donnell C 2012; Ligi I et al. 2008). The catastrophic impact of medication errors is demonstrated by numerous high-profile cases in the UK such as Richie William's death from inadvertent intrathecal rather than intravenous administration of vincristine, for which two doctors faced charges of manslaughter (Dyer C 1999).

The student's recent publication reported paediatric safety incidents from general practice and showed that vaccination-related errors were among the most frequently reported and harmful safety incidents affecting children (Rees P et al. 2015). These errors included inadvertent administration of the wrong vaccine, the wrong number of vaccine doses, and administration at the wrong time. The sometimes-catastrophic effect of deviating from the vaccination schedule has been demonstrated by child deaths from vaccine-preventable infections and recent Measles outbreaks in the UK and USA (Greaves F and Donaldson L 2013; Harnden A et al. 2009).

Child death reviews highlight safety incidents in primary care such as: failure to recognise and manage severe infection, failure to vaccinate, and failure to follow up patients. Those with chronic conditions contribute substantially to UK child deaths (Harnden A et al. 2009). Thomson MJ et al. 2006 highlight failures in first-access services within primary care by reporting that 50% of children with diagnosed meningococcal infection had been sent home from their first consultation, delaying diagnosis of a life-threatening infection.

The litigation costs of missed diagnoses in children equates to over £20million in the past 13 years (Wolfe I et al. 2011). These safety incidents underpin the poor UK performance in numerous measures of paediatric care quality, as highlighted in section 1.1.

Approximately 6000 children die each year in England and Wales, 20% of these deaths are thought potentially preventable with better care, however it is unclear how many of these deaths were the result of a patient safety incident (Pearson G 2008; Sidebotham P et al. 2014b).

1.3 Methods of measuring the burden and assessing the nature of unsafe care and healthcare error

Efforts to improve care quality to date have been hindered by unreliable data on the prevalence, burden and nature of substandard care. However, before paediatric care quality can be addressed and improved, widespread and reliable measurement of the scale of the problem, including the most frequent and harmful sources of substandard care, is required (Walsh KE et al. 2014). These measurements must be both accurate and precise (Thomas EJ and Petersen LA 2003). Several methods exist for assessing care quality; key methods will be presented and their respective attributes and weaknesses discussed.

1.3.1 Definitions

Numerous terms exist to describe the outcomes of unsafe care and different methods focus on different outcomes such as: preventable harms, all harms, and near misses.

Preventable harm is defined as a negative patient outcome "that would not have occurred if the patient had received ordinary standards of care appropriate for the time" (World Health Organisation 2009). Harm, preventable or unpreventable, is defined as a negative patient outcome that includes "temporary or permanent impairment of the physical, emotional, or psychological function or structure of the body and/or pain requiring intervention" (World Health Organisation 2009).

Near misses however also encompass errors that do not result in harm and are defined as "events or situations that could have resulted in an accident, injury or illness, but did not, either by chance or through timely intervention" (World Health Organisation 2009). Near misses represent an important group of errors because the failures and system weaknesses underpinning them tend to be the same as those contributing to harmful errors. Unfortunately near misses can be difficult to detect and are often overlooked by certain measurement methods. Similarly some methods focus solely on preventable harm whereas others focus on all harm arguing that what constitutes preventable harm is continuously evolving (Parry G et al. 2012)

1.3.2 National-level administrative data

The Royal College of Paediatrics and Child Health (RCPCH) conducted a retrospective epidemiological review of all-cause mortality in UK children, using nationally representative and coded longitudinal administrative data linked to supplementary data sources such as birth and death certificates (Hardelid P et al. 2013). These datasets are likely to be linked to national-level longitudinal primary care data, mental health services data, and data on emergency department attendances and intensive care stays in the near future (Dattani N et al. 2013; Hardelid P et al. 2013). There has been no national-level measurement of paediatric care quality in the UK. However, these data could be monitored for poor quality care using pre-specified care quality indicators and their respective ICD-10 codes, to estimate the burden of substandard care, as exemplified in the USA by the Harvard practice medical study (Brennan TA et al. 1991; Brennan TA et al. 2004; World Health Organisation 2010).

There are numerous benefits to using nationally collected longitudinal administrative data. However, utilising these data effectively is expensive and time consuming; and it is widely acknowledged that the data are too crude to provide meaningful information about the complexities of care. Despite the availability of national epidemiologic data about child health, their value is limited by the quality of available data and the systems providing data (Fraser J et al. 2014). For example, a Confidential Enquiry highlighted that 35% of child death certificates had an incorrect cause of death recorded (Fraser J et al. 2014; Wolfe I et al. 2014). Considerable delay in death registration and certification combined with inaccurate and incomplete records limit the utility and depth of learning that can be gained from this approach. However, the RCPCH advocates supplementing data linkage of routinely collected data with in-depth case note reviews to address these criticisms (Hardelid P et al. 2013).

1.3.3 Case note reviews

Historically patient medical records have been used to measure the incidence and burden of substandard care (Parry G et al. 2012). A random sample of medical records are reviewed using pre-defined 'quality indicators' to identify the proportion of children receiving appropriate care, and the proportion receiving substandard care. This approach provides case note reviewers with the opportunity to identify detailed clinical information to understand and contextualise the care quality incident. Traditionally this method has

been used to estimate the incidence of substandard care and to investigate the types of failures leading to poor care quality (Mangione-Smith R et al. 2007b;Sari ABA et al. 2007;Vincent C et al. 2001). Gill P et al. 2014 have developed quality indicators for children in UK family practice, however they have not yet been tested. The extent of substandard care can be monitored temporally and the impact of improvement initiatives assessed with this approach, locally and nationally.

Reviews may be disease focused e.g. care quality in children with asthma, or they may be restricted to serious cases or specific child deaths. Disease-based reviews of care quality in live children are also becoming increasingly popular and allow crude comparisons of care quality with other countries (Asthma UK 2007; Wolfe I et al. 2014). They are conducted annually in the UK for diabetic care and the RCPCH have recently finished a three-year audit of epilepsy care quality (Royal College of Paediatrics and Child Health 2012).

Reviews of medical records are expensive, time-consuming, labour-intensive, inquisitorial and prone to incomplete ascertainment (Mangione-Smith R et al. 2007b; Parry G et al. 2012; Sari ABA et al. 2007; Vincent C et al. 2001). For example, this method relies on the content and the quality of medical records, which are often subjective or incomplete. Hardelid P et al. 2013 have demonstrated that the issue of incomplete medical records or ascertainment can be overcome-at least partially-by linking data sources.

1.3.3.1 Trigger tools

Trigger tools offer an alternative and more efficient approach to reviewing medical records compared with traditional methods of systematically reviewing complete records. This method of harm detection and measurement includes selecting a random sample of medical records for case note review (Chapman SM et al. 2014). These records are then systematically searched, typically by a trained professional, for 'triggers' which are a predefined list of events that suggest patient harm such as hypoglycaemia (Chapman SM et al. 2014). Records where triggers are identified are reviewed in-depth to determine whether the trigger represents an adverse event in that patient. When adverse events and harm are suspected, second reviewers that are typically clinicians, confirm the occurrence of harm and its severity (Chapman SM et al. 2014).

Tools such as the Global Trigger Tool have been tested and developed for adult care in specific care settings including: acute hospitals, surgery, critical care, and primary care (Classen DC et al. 2011;Griffin FA and Classen DC 2008;Griffin FA and Resar RK 2009;Parry G et al. 2012;Resar RK et al. 2006;Singh R et al. 2009). Fewer paediatric-specific trigger tools exist, however they have been developed for hospital setting (Agarwal S et al. 2010;Chapman SM et al. 2014;Larsen GY et al. 2007;Matlow AG et al. 2011;Muething SE et al. 2010;Sharek PJ et al. 2006;Takata GS et al. 2008). A UK based paediatric trigger tool for use in hospitals was only recently developed and to the student's knowledge no primary care specific paediatric trigger tools exist (Chapman SM et al. 2014).

This method is more sensitive than comparable methods since it detects more adverse events (Parry G et al. 2012). Several factors may underlie this sensitivity: trigger tools focus on harm rather than error, and they do not rely on recognition and reporting of error or harm, they are therefore more likely to detect errors of omission (Chapman SM et al. 2014; Parry G et al. 2012). Additional attributes include the low cost associated with the use of this tool to obtain relevant data for improvement, and the ability of this method to provide reliable data, making it suitable for learning at a local level. Studies using trigger tools have shown high inter-rater reliability between reviewers (Chapman SM et al. 2014). However the validity of estimates of harm measurements generated by trigger tools is dependent on the accuracy and completeness of documentation.

The application of trigger tools is currently labour intensive although electronic medication records could support the automated detection of triggers in the future (Chapman SM et al. 2014; Parry G et al. 2012; Walsh KE et al. 2014). Despite the high reliability and harm detection rate of trigger tools, they were designed to complement other methods of harm detection rather than replace them (Chapman SM et al. 2014; Parry G et al. 2012; Walsh KE et al. 2014).

1.3.4 Incident reporting systems

Reporting systems are a well-established resource widely used in healthcare to provide insights into unsafe and poor quality care (Rees P et al. 2015). The purpose of an incident reporting system is to enable an organisation and its staff to learn from human and system errors to prevent their reoccurrence (Vincent C 2007; World Alliance for Patient Safety 2005). A successful reporting system relies on staff submitting reports, good quality descriptions, high quality analysis of reports, and responding to those findings to improve

safety (World Alliance for Patient Safety 2005). High-risk organisations such as those in the nuclear, petrochemical, and aviation industry value the contribution of safety reports (World Alliance for Patient Safety 2005). They facilitate identification of system weaknesses that require remedy to prevent disaster. The healthcare industry has followed the lead of such industries in their approach to safety, including the development of incident reporting systems.

Incident reporting systems vary in terms of: their purpose, who reports, what is reported, how to report, the analysis of reports, and the dissemination of findings (World Alliance for Patient Safety 2005). Reports tend to come from healthcare professionals although there is increasing patient participation in this field and some reporting systems are open to receiving reports from patients, families and patient advocate groups. Some reporting systems only receive reports of adverse drug reactions, for example, the yellow card system in the UK, whereas other systems receive reports of any adverse event such as the national reporting system in Denmark (World Alliance for Patient Safety 2005). These reports are analysed using various classifications and some reports may prompt more detailed investigations such as a root cause analysis (National Patient Safety Agency). The findings of analysis, including recommendations for improvement are then disseminated to stakeholders such as reporters.

Reporting systems can be used to inform improvements at the national or local level. At the national level, aggregating large quantities of data enables detection of rare incidents that would otherwise be missed (Rees P et al. 2014). For example, the UK National Reporting and Learning System (NRLS) was integral in detecting the association of bone cement implantation syndrome and the use of cement in hip fracture surgery (Panesar SS et al. 2009a; Panesar SS et al. 2009b). However hazards particular to a hospital or ward can also be detected by analysing reports submitted locally, for example, to identify outdated or faulty equipment. Reporting can result in improvement. For example, through circulating alerts about significant new hazards such as drug side effects; by disseminating learning from serious incidents not only to the institution where the incident occurred but to other institutions; and by generating recommendations for best practice to mitigate identified hazards and system failures (World Alliance for Patient Safety 2005).

Incident report data have successfully informed over 300 studies in the published literature. Such research has identified problem areas in secondary care requiring improvement including: reliable insulin administration, early detection of surgical

complications, and lithium prescribing and monitoring, highlighting the potential value of incident report data (Cresswell KM and Sheikh A 2008; Lamont T et al. 2011a; Lamont T et al. 2011b; Lamont T et al. 2010; Lamont T et al. 2011c).

Despite their unique advantages, such as being well placed to detect near misses, incident reporting systems have widely published weaknesses. The Achilles heel of incident reporting systems is their inability to provide a complete picture of healthcare safety due to under-reporting (Vincent C 2007; Vincent C et al. 2008; Vincent CA 2004). A case note review concluded that only 7% of detected incidents are reported to the UK reporting system, and that those incidents reported are subject to numerous biases (Sari ABA et al. 2007). Variability in the content of reports and what gets reported between different professional groups, wards, and organisations creates 'reporting bias', which is a considerable challenge for those seeking to interpret meaning and generate learning from reports.

The Berwick report emphasises the importance of a well-designed reporting system within every healthcare organisation (Department of Health 2013). The aim of such a system is not to capture all incidents, or to capture them representatively, but to provide a window into the hazards and system failures that impact patient care. Many claim that we are drowning in big data. The challenge is to effectively utilise data that are routinely collected, such as incident report data. Well publicised healthcare failures such as those in Mid-Staffordshire and the Bristol Heart Inquiry would not necessarily have been prevented with increased data collection (Francis R 2013;Kennedy I 2001). Numerous reports and complaints of unsafe care were made at these institutions and sufficient data were collected by the organisations to demonstrate worrying trends in mortality, yet these trends went unacknowledged (Panesar SS et al. 2013b).

The WHO is developing a minimal information model for patient safety reporting, which will allow international comparison of incident report data (World Health Organisation). The limitations of incident reporting systems are well known and perhaps over-emphasised as they provide vast quantities of data for learning (Carson-Stevens A et al. 2015). Many believe that incident report data are under-valued and that they could be better utilised globally to provide insights into the safety of healthcare, and to facilitate improvements in care quality.

1.4 National Reporting & Learning System (NRLS)

Patient safety incident reporting systems provide one lens through which to view human and system failures that may result in harm to patients. This is the purpose of the NRLS which collects reports of any "unintended or unexpected incident that could have or did lead to harm for one or more patients receiving NHS funded care" in England or Wales, to identify risks at a national level(National Reporting and Learning System).

The NRLS was established in 2003 in response to a governmental report 'An Organisation with a Memory', which criticised the ability of the NHS to identify and address serious failures in healthcare at that time (Department of Health 2000). It was managed by the National Patient Safety Agency (NPSA) - an independent body established in 2000 whose purpose was to "implement and operate the [NRLS] - to improve patient safety by reducing the risk of harm through error" (Department of Health 2001). However, control of the NRLS was transferred to the NHS commissioning board in 2012.

Despite calls for mandatory reporting of adverse events and specific near misses, at its creation the NRLS was completely voluntary and anonymous. The intention was to minimise defensive reporting and disincentives to reporting, such as blame or fear of retribution encouraging full and candid reporting (Department of Health 2000; Francis R 2013). In addition, the NPSA did not have the power to enforce mandatory reporting and the purpose of reporting was to galvanise learning rather than to detect and compare failures between healthcare organisations. However, in 2010 reporting of incidents resulting in severe harm or patient death became mandatory (Francis R 2013).

Staff and patients are encouraged to report and this can be done directly and independently to the NRLS online. Staff can also report incidents within their place of work using local reporting procedures, and in specialties such as anaesthesia they can complete specialty-specific reports (Francis R 2013). All reports submitted locally are analysed by that parent healthcare organisation for 'local learning' and then all reports are anonymised and uploaded to the NRLS by a designated person for 'national learning' (see Figure 1) (Donaldson LJ et al. 2014; Rees P et al. 2015).

Every report form contains 'essential information' such as patient age, but the layout and appearances of report forms vary between organisations. For example, some include instructions to reporters about what information to include and others do not. NRLS reports contain multiple categorical variables that include patient age range, location,

care setting, country, and severity of harm. They also contain free-text information about what happened, potential causes, and suggestions for future prevention (Donaldson LJ et al. 2014; Rees P et al. 2015).

The NHS commissioning board cannot scrutinise every report submitted to the NRLS; it receives over 65,000 paediatric related reports per annum, despite considerable underreporting (Carson-Stevens A et al. 2015).-The detail and accuracy of information included in reports is at the discretion of reporters, who often report with hindsight, and are prone to inherent human biases (Donaldson LJ et al. 2014;Rees P et al. 2015). Despite these limitations, the NRLS has a well-established infrastructure, and receives large quantities of rich data that can be used for learning and hypothesis generation (Rees P et al. 2015).

The insights offered by incident report data and the potential benefits of its analysis are clear despite its limitations, yet no systematic analysis of primary care-related paediatric safety incidents submitted to the NRLS has been conducted to date.

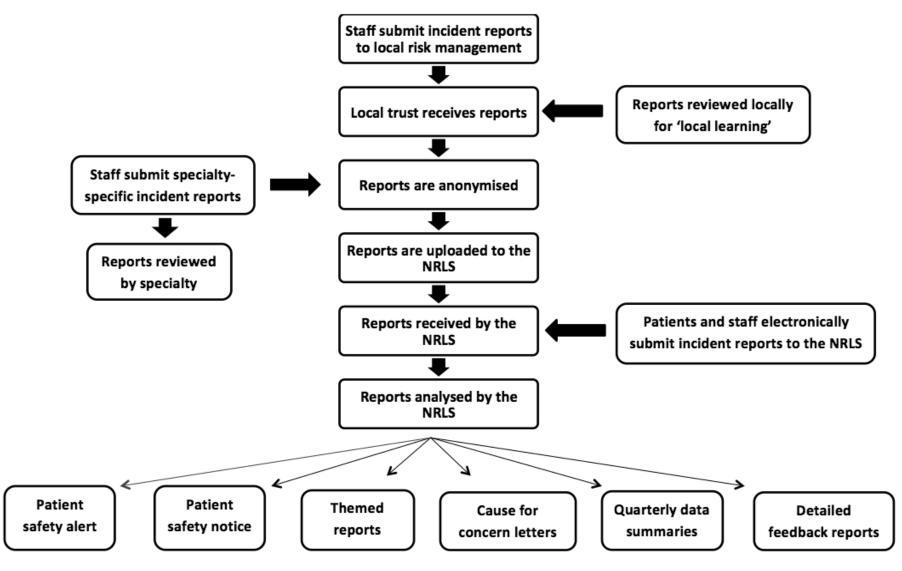


Figure 1 page 21: flow diagram illustrating the process of reporting incidents to the NRLS and how the NRLS responds to reports

1.5 Aims and objectives

1.5.1 Aim

To explore the nature, contributory factors, severity, and outcomes of paediatric safety incident reports related to vaccination in primary care

1.5.1.1 Objectives

- Provide an overview of the data included and processed, and present key quantitative findings
- Present the key themes and sub-themes identified from thematic analysis of purposively sampled reports, and describe how these themes relate to each other
- Combine the insights gained from the quantitative and qualitative analysis, and explain how the qualitative insights relate to the quantitative findings

1.5.2 Aim

To explore the nature, contributory factors, severity, and outcomes of paediatric safety incident reports involving 'unwell' children in primary care

1.5.2.1 Objectives

- Provide an overview of the data included and processed, and present key quantitative findings
- Present the key themes and sub-themes identified from thematic analysis of purposively sampled reports, and describe how these themes relate to each other
- Explain how the qualitative insights related to the quantitative findings

1.5.3 Aim

To conduct a literature review to identify quality improvement interventions tested, or implemented to improve either the paediatric vaccination process or care for 'unwell' children.

Chapter 2: Method

This chapter will describe how the student decided to utilise the NRLS data to best achieve the aims and objectives previously described. The methods used to sample, search, process, and analyse the NRLS data will be presented, whilst justifying how they were chosen and comparing them to alternative methods. The methods used to conduct the literature review will be presented alongside the literature review in section 3.3, as the review question and inclusion criteria are dependent on the results presented in sections 3.1 and 3.2.

2.1 Data handling

2.1.1 Data sampling

Of 272,884 primary care-related incident reports, 20,118 reports involving children were identified on applying a filter to the patient age-related columns in Microsoft excel. These reports were searched using the search strategies.

2.1.2 Search strategy

The student designed two search strategies for maximal recall of vaccination-related incidents and incidents related to 'unwell' children (Appendix 2.1 and 2.2). Previously analysed reports were reviewed to extract key terms and alternative spellings/misspellings of these terms pertinent to the above topics. This list of vaccination-related key terms derived from the reports themselves was combined with lists of vaccine brands and vaccine generic names from the children's British National Formulary (BNF) (Paediatric Formulary Committee 2013). Similarly the search strategy to identify reports of 'unwell' children was iterated using appropriate terms from the International Classification of Disease 10 (ICD-10) (World Health Organisation 2010).

Both search strategies were reviewed and adapted by clinicians (the student's supervisors). All paediatric reports were imported into a qualitative software package (NVIVO 9, QSR International), where the key terms were used to search the reports. Alternative spellings, synonyms and abbreviations used in the NRLS reports were identified using NVIVO text search functions and these additional terms were incorporated into the search strategy to maximise its sensitivity.

2.1.3 Data processing

Once the relevant reports were identified the student imported them into a separate password protected Microsoft Excel document. The free text components of reports were read and the incident types, potential contributory factors, severity of harm and incident outcomes were classified. The vaccines involved in the vaccination-related incidents were noted. The pre-existing and/ or presenting diagnosis signs or symptoms of 'unwell' children were classified using ICD-10; and the medications involved in incidents were classified using the children's BNF. The inclusion and exclusion criteria are listed in Table 1. During data processing potentially severe near misses, and rare or theoretically important reports were logged for subsequent qualitative analysis.

Table 1 page 25: inclusion and exclusion criteria

Inclusion	Exclusion
Incidents in children aged <18 years	Reports without a pre-allocated age group
AND Incidents occurring in primary care	Incidents occurring in secondary care but reported by primary care
AND Incidents occurring between 2002-2013	Incidents involving child maltreatment,
	sexual abuse, fabricated or induced illness,
	emotional abuse, or neglect
AND Vaccination-related incidents OR	Incidents without free-text descriptions
incidents involving 'unwell' children	
defined as children with any described	
symptoms, signs, diagnoses, or prescribed	
medications implying illness	

2.1.4 Classification frameworks

In order to comprehensively describe the detailed free-text descriptions contained within reports, the student sought to classify incident types, contributory factors, incident outcomes, and severity of harm. Existing classifications were reviewed for suitability (Computer Sciences Corporation 2015;Dovey SM et al. 2002;Jacobs S et al. 2007;Kostopoulou O and Delaney B 2007;Makeham MA et al. 2002;Makeham MA et al. 2008;Rubin G et al. 2003;The Netherlands: Eindhoven University of Technology;West D et al. 2005;World Health Organisation 2009).

I had originally intended to use the Learning from International Networks about Errors and Understanding Safety in Primary Care [LINNAEUS Euro-PC] taxonomy to classify reports. This taxonomy was designed for use in primary care to classify incident type and had been used in previous studies (Martijn LLM et al. 2013;Rosser W et al. 2005;Woolf SH et al. 2004). During the student's previous work with NRLS data, the incident type and contributory factors frameworks were separated into two parent categories: system factors and human factors mirroring the LINNEAUS taxonomy, and contributory incidents were in both the incident descriptor framework and contributory factors framework (Rees P et al. 2015). These frameworks were not 'user friendly', had extensive overlap, and codes within frameworks were not mutually exclusive.

The student re-organised the incident type and contributory factor frameworks, and this re-organisation was influenced by the WHO International Classification for Patient Safety (World Health Organisation 2009). The amended incident descriptor framework included incidents and contributory incidents, aggregated under ten parent codes: administration, documentation, referral, diagnosis and assessment, treatment and procedures, medications and vaccines, investigations, communication, care equipment, and 'other'. The system and human factors e.g. referral process and referral decision, remained although would now have the same parent code e.g. referral; this facilitated the process of logically classifying incidents.

The 'new' contributory factors framework included contributory factors only, and not contributory incidents, as it had previously. These contributory factors were re-organised under four parent categories: patient/caregiver factors, staff factors, equipment/vaccine/ medication factors, and organisational factors, mirroring Eindhoven's

Classification and WHO International Classification for Patient Safety (The Netherlands: Eindhoven University of Technology; World Health Organisation 2009).

Previous studies had used the ICD-10 to classify incident outcomes, however the student decided that the ICD-10 was too extensive for use with this dataset (World Health Organisation 2010). Conversely during a previous study, the student used Vincent's typology of harm to classify the type of harm outcomes resulting from incidents and found this framework too broad (e.g. there were only 6 types of harm) (Vincent C et al. 2013). Therefore, an in-house framework grounded in UK NRLS data was created.

Therefore, three of the four frameworks used were developed 'in-house', grounded in UK primary care data, and iterated extensively to accurately capture the descriptions of incidents, contributory factors and outcomes within reports.

Prior to deciding on a framework to classify harm severity the student reviewed three of the most frequently used frameworks: the NRLS classification, severity assessment code matrix, and WHO International Classification for Patient Safety (National Reporting and Learning System; New South Wales Government 2005; World Health Organisation 2009). Tables 3-5 demonstrate the definitions of harm severity used in each classification: there is little variation in the definitions of harm at the patient level but the severity assessment code matrix classifies harm more comprehensively.

The severity assessment code matrix was piloted on small sample of reports to classify the severity of harm at the patient, staff, services, financial, and environmental levels where free-text descriptions allowed (New South Wales Government 2005). However, as reports only described harm at the patient level the student decided that such a comprehensive classification matrix was unnecessary and incompatible with this dataset.

The student decided to classify harm severity using the WHO International Classification for Patient Safety as the categories mirror those of the NRLS (no harm, low harm, moderate harm, severe harm, death) but the definitions are more comprehensive (National Reporting and Learning System; World Health Organisation 2009). This will also aid comparison with previous and future studies of this nature.

All reports submitted to the NRLS contain a harm severity decided by the reporter at the originating healthcare organisation. The reporter-allocated harm severity was only used

during data processing when there was insufficient detail in the free-text descriptions to re-classify the severity of harm using the WHO International Classification for Patient Safety. Thus, the severity of harm was upgraded or downgraded from the reporter-allocated harm severity when additional information was available, and the free-text descriptions of harm severity differed to the reporter-allocated harm severity.

Table 2 page 29: the severity assessment code matrix (New South Wales Government 2005)

Severity of harm	Minimum	Minor	Moderate	Major	Serious
Patient level	Patients with No injury or increased level of care or length of stay	Patients requiring Increased level of care including: Review and evaluation Additional investigations Referral to another clinician	Patients with permanent reduction in bodily functioning (sensory, motor, physiological, or psychological) unrelated to the natural course of the illness and differing from the expected outcome of patient management or any of the following: Increased length of stay as a result of the incident Surgical intervention required as a result of the incident	Patients suffering a major permanent loss of function (sensory, motor, physiological) unrelated to the natural course of the illness and differing from the expected outcome of patient management or any of the following: Suffering significant disfigurement as a result of the incident Patient at significant risk due to being absent against medical advice Threatened or actual physical or verbal assault of patient requiring external or police intervention	Patients with death unrelated to the natural course of the illness and differing from the immediate expected outcome of the patient management or: Suspected suicide Suspected homicide Or any of the following: National sentinel events Procedures involving the wrong patient or body part Suspected suicide in hospital Retained instruments Unintended material requiring surgical removal Medication error involving the death of a patient Intravascular gas embolism Haemolytic blood transfusion Maternal death associated with labour and delivery Infant discharged to the wrong family
Staff level	No injury or review required	First aid treatment only with no lost time or restricted duties	Medical expenses, lost time or restricted duties or injury / illness for 1 or more staff	Permanent injury to staff member, hospitalisation of 2 staff, or lost time or restricted duty or illness for 2 or more staff or pending or actual WorkCover prosecution, or threatened or actual physical or verbal assault of staff requiring external or police intervention	Death of staff member related to work incident or suicide, or hospitalisation of 3 or more staff
Visitors level	No treatment required or refused treatment	Evaluation and treatment with no expenses	Medical expenses incurred or treatment of up to 2 visitors not requiring hospitalisation	Hospitalisation of up to 2 visitors related to the incident / injury or pending or actual WorkCover prosecution	Death of visitor or hospitalisation of 3 or more visitors
Services level	Services: No loss of service	Reduced efficiency or disruption to agency working	Disruption to users due to agency problems	Major loss of agency / service to users	Complete loss of service or output
Financial level	No financial loss	Loss of assets replacement value due to damage, fire etc. to \$50K	Loss of assets replacement value due to damage, fire etc. \$50K to \$100K or loss of cash/investments/assets due to fraud, overpayment or theft to \$10K	Loss of assets replacement value due to damage, fire etc. \$100K-\$1M, loss of cash/investments/assets due to fraud, overpayment or theft \$10K-\$100K or WorkCover claims \$50K-\$100K	Loss of assets replacement value due to damage, fire etc. > \$1M, loss of cash/investments/assets due to fraud, overpayment or theft >\$100K or WorkCover claims > \$100K
Environmental level	Nuisance releases	Off-site release contained without outside assistance	Off-site release contained with outside assistance or fire incipient stage or less	Off-site release with no detrimental effects or fire that grows larger than an incipient stage	Toxic release off-site with detrimental effect. Fire requiring evacuation

Table 3 page 30: National Reporting & Learning System definitions of harm severity (National Reporting and Learning System)

Severity of harm	No harm	Low harm	Moderate harm	Severe harm	Death
Definitions	Impact prevented - any	Any patient safety	Any patient safety incident	Any patient safety	Any patient safety
	patient safety incident	incident that required	that resulted in a moderate	incident that appears to	incident that directly
	that had the potential to	extra observation or	increase in treatment and	have resulted in	resulted in the death of
	cause harm but was	minor treatment and	which caused significant but	permanent harm to one	one or more persons
	prevented, resulting in no	caused minimal harm,	not permanent harm, to one	or more persons	receiving NHS-funded
	harm to people receiving	to one or more	or more persons receiving	receiving NHS-funded	care
	NHS-funded care	persons receiving NHS-	NHS-funded care	care	
		funded care			

Table 4 page 30: WHO International Classification for Patient Safety definitions of harm severity (World Health Organisation 2009)

Severity of harm	No harm	Mild harm	Moderate harm	Severe harm	Death
Definitions	Patient outcome is not	Patient outcome is	Patient outcome is	Patient outcome is	On balance of
	symptomatic or no	symptomatic, symptoms	symptomatic, requiring	symptomatic, requiring	probabilities, death was
	symptoms detected and	are mild, loss of function	intervention (e.g.,	life-saving intervention	caused or brought
	no treatment is required	or harm is minimal or	additional operative	or major	forward in the short
		intermediate but short	procedure; additional	surgical/medical	term by the incident
		term, and no or minimal	therapeutic treatment),	intervention, shortening	
		intervention (e.g., extra	an increased length of	life expectancy or	
		observation, investigation,	stay, or causing	causing major	
		review or minor treatment)	permanent or long term	permanent or long term	
		is required	harm or loss of function	harm or loss of function	

2.1.5 Models of incident analysis

Patient safety incidents are typically complex with multiple factors preceding them. The free text in each incident report can provide sufficient detail to identify the related incidents and contributory factors. Numerous studies have summarised free-text descriptions of these complex events by selecting only one incident type, predominantly the most important or severe, the first incident occurring in the chain of causality, or the final incident in the chain of causality (Donaldson LJ et al. 2014;Dovey SM et al. 2002;Magrabi F et al. 2011;Panesar SS et al. 2013a).

The student did not want to reduce or summarise the rich free-text descriptions within reports and sought to capture the complex relationships described within them. Other studies aiming to do this categorised multiple incidents and contributory factors, and some did this chronologically to model the chain of causality (Rees P et al. 2015;Suresh G et al. 2004;Woolf SH et al. 2004). However this approach gave rise to confusion between incidents-particularly contributory incidents-and contributory factors i.e. there was significant overlap between them (Hibbert PD et al. 2007;Rees P et al. 2015;Suresh G et al. 2004).

To address the above issues the student decided to use the Australian Recursive Model of Incident Analysis to organise and order data processing (Hibbert PD et al. 2007). This facilitates consistency in the application of codes and allows in-depth classification of reports. Additionally, unlike other models, the recursive model has clear rules differentiating primary incidents, contributory incidents, and contributory factors (Hibbert PD et al. 2007). This approach was therefore compatible with the frameworks developed in-house i.e. there was no overlap between the incident descriptor framework and the contributory factor framework.

There are nine incident analysis rules in this model: the primary incident type is the incident chronologically proximal to the patient prior to harm/outcome (or potential harm) (Table 5) (Hibbert PD et al. 2007). Therefore, by definition, the outcome of the primary incident type cannot be an incident it can only be an outcome. Incidents that occur prior to the primary incident type (more distal to the patient) that played a part in the occurrence of the primary incident are by definition contributory incidents. Contributory factors cannot be incidents in their own right, such as a child being looked-after; these are found in the contributory factors framework. These sequential incidents

are coded in reverse chronological order, which is similar to the approach used in a root cause analysis. (Figures 2 and 3) Working backwards from the primary incident type and mapping out the sequence of incidents, contributory incidents, and contributory factors, allows us to reach the root cause (where there is sufficient detail).

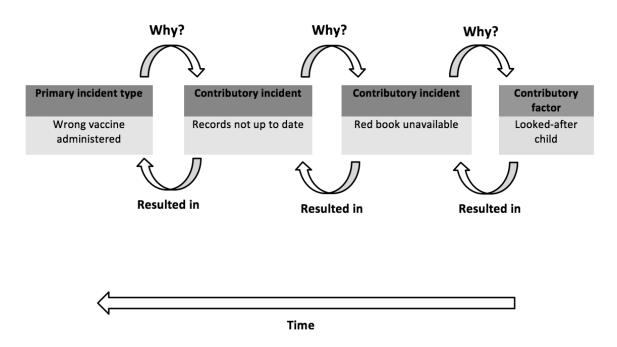


Figure 2 page 33: illustrates the Australian Recursive Model of Incident Analysis

		Incident Types				Number of incidents	Contrib	utory Factors	Number of contributory factors
Wrong vaccine administered	6.11.3.1	Records not up to date	2.1.3	Red book unavailable	2.1.1.1		3 Looked-after child	1.5	1

Figure 3 page 33: an example of the layout of the Recursive Model of Incident Analysis during data processing in Microsoft Excel.

Table 5 page 34: the nine rules of the Australian Recursive Model of Incident Analysis (Hibbert PD et al. 2007; Rees P et al. 2015)

Incident Analysis Rules	Rule Example
1. An incident has a set of	Missed diagnosis (incident) because the physician
contributory factors and / or	did not adequately examine the patient
contributory incidents	(contributory incident) and the physician had
	inadequate knowledge (contributory factor)
2. An incident can contribute to	Missed diagnosis (contributory incident) resulted in
another incident	a patient not receiving a timely referral to the
	hospital (primary incident)
3. Contributory factors cannot be	A mistake (contributory factor <i>not</i> an incident)
incidents in their own right	resulting in the wrong prescribed medication dose
	(primary incident)
4. An incident has a set of	Wrong prescribed medication dose (primary
outcomes	incident) resulting in a medication overdose and
	hospital admission (outcomes)
5. An incident can be an outcome	Records not up to date (contributory incident)
of another incident	resulting in the wrong prescribed medication
	(primary incident and outcome)
6. Some outcomes cannot be	Admission to hospital (outcome) following the wrong
incidents in their own right	prescribed medication (primary incident)
7. An outcome of an incident	Communication incident between care providers
could be a contributory incident	(contributory incident) resulting in records not
to another incident	being up to date (contributory incident and
	outcome), resulting in a referral incident (primary
	incident)
8. An incident can be designated	Communication incident (incident) leading to
the primary incident type - the	inaccurate records (incident), leading to the wrong
incident proximal to the	prescribed medication (primary incident type)
descriptive patient outcome	
9. The outcome of a primary	Admission to hospital (outcome) following the wrong
incident cannot be an incident	prescribed medication dose (primary incident type)

2.1.6 Quantitative analysis

Prior to analysis, the types of conditions present in 'unwell' children classified using the ICD-10 were aggregated into thirteen broad categories (Table 6).

The processed data underwent descriptive analysis to illustrate the frequency distribution of variables including: age, time, incident types, contributory factors, severity of harms, outcomes, types of conditions and medications or vaccines implicated in incidents. The relationships between these variables were explored using numerous cross-tabulations for example: incident type by age, incident type by contributory factors, primary incident type by contributory incidents, contributory factor one by contributory factor two, incident type by diagnosis, incident type by harm, incident type by outcome *etc*. This provided descriptions of temporal trends, highlighted potential associations between variables and combinations of contributory factors.

The relationship between the primary incident type and contributory incidents, primary incident type and contributory factors, and various contributory factors with each other were examined to identify clusters of factors.

Table 6 page 36: illustrates how the ICD-10 codes for children's pre-existing and/ or presenting conditions were grouped

Type of condition	ICD 10 codes
Infections	A00-99; B00-99
Cancer and blood	C00-97; D00-89; R70-79
Skin and musculoskeletal system	L00-99; M00-99; Q65-79; R20-23; R25-29
Neurological and sensory system	G00-99; H00-95; Q00-Q07; Q10-18
Respiratory system	J00-99; Q30-34; R04-6; R09.0-3
Mental and behaviour	F00-99; X60-84
Injuries	S00-99; T00-98; V01-X59; X85-Y09; Y10-Y34; Y35-36; Y40-84; Y85-89
Circulatory system	Q20-28; R00-03; R09.8
Digestive and genitourinary system	K00-93; N00-99; Q35-37; Q38-45; Q50-56; Q60-64; R10-19; R30-39; R80-82
Endocrine, metabolic and nutrition	E00-90
Non-specific signs and symptoms	R07; R40-46; R47-49; R50-69; R83-89; R90-94; R95-99
Pregnancy, chromosomal other congenital disorders	O00-99; P00-96; Q80-89; Q90-99

2.2 Thematic analysis

2.2.1 Justification of qualitative method

Thematising data features in numerous types of qualitative methods such as grounded theory and ethnography and should therefore be considered a fundamental tool of qualitative analysis (Holloway I and Todres L 2003); (Braun V and Clarke V 2006). Despite this overlap Braun V and Clarke V 2006 argue that it should also be considered a standalone method.

An attribute of thematic analysis is its flexibility: it is not constrained to a particular epistemology or ontology. The student recognised that it was important to take a realist approach and capture the content of reports i.e. what reporters wrote. However, the student also recognised the importance of seeking the underlying meaning of the reports' content i.e. what the reporters meant or implied, which is consistent with an idealist approach (Bourgeault I et al. 2010; Denzin NK and Lincoln YS 2011). Thematic analysis offered the flexibility to use both ontological approaches, realist and idealist, where the data allowed, to augment learning (Braun V and Clarke V 2006).

Braun V and V Clarke 2006 describe two thematic analysis methodologies: theoretical and inductive. A mixture of these was used in this study. Analysis was grounded in the data (data-driven) in an inductive manner. New theoretical insights about the human and system failures underpinning incidents were noted and sought, and there was no pre-defined framework. However, Braun V and Clarke V 2006 argue that data cannot be coded in an epistemological vacuum and prior data processing and clinical knowledge did inform the student's theoretical insights, which in turn influenced sampling. These theoretical insights were further explored through thematic analysis of the purposively sampled reports.

2.2.2 Qualitative sampling

Reports that provided new insights, and supported or contradicted emerging theories during classification, were selected for qualitative analysis, providing a purposive sample of reports.

2.2.3 Qualitative analysis

Purposively sampled reports were imported into NVIVO 9 (QSR International), and read and re-read for familiarisation with the data. Systematic searching of the data was conducted to identify deviant or noteworthy cases. Broad brush coding using word frequency queries in NVIVO (also known as content analysis) was conducted for orientation to the data. After immersion in the data, segments of free-text underwent open coding in an inclusive descriptive manner, line by line, to capture every nuance and contextual factor, to preserve the reports' meaning and not summarise or reduce the data ((Bryman A 2012; Glaser BG and Strauss AL 2009; Miles MB and Huberman AM 1994; Pope C and Mays N 1995; Pope C et al. 2000). Nodes were created in NVIVO for this purpose, which often incorporated key words or phrases used in the reports (in-vivo codes) to preserve reporters' views. Often multiple nodes were created for the same section of text where the description satisfied inclusion into multiple nodes. Semantic (descriptive) features and latent (interpretative) features were coded. A constant comparative approach was used, this involved comparing each node with previously coded text to develop analytic categories, an iterative process (Denzin NK and Lincoln YS 2011; Lacey A and Luff D 2001).

Descriptive nodes were extracted and underwent analysis at the latent and interpretive level, where the reports were re-examined to extract underlying meaning and inference. Overlapping nodes were then merged into subthemes that created an index system (Bazeley P 2007). Free-text was re-coded using this index system. Sub-themes were refined, and combined to form over arching themes. The relationships between these sub-themes and overarching themes were mapped, analysed and interpreted (Pope C et al. 2000). Themes were not determined by the prevalence of a concept but its perceived clinical and theoretical importance. Themes and the hierarchy of sub-themes within them were examined, adapted, and refined to ensure internal homogeneity and external heterogeneity. Where necessary, data were re-coded for consistency and comprehensiveness: typical of the organic process of qualitative analysis (Braun V and Clarke V 2006).

Theoretical insights were concurrently documented to create an audit trail and aid reflexivity (Bazeley P 2007).

2.3 Mixed methods synthesis of results

A mixed method is defined as "integrating quantitative and qualitative data collection and analysis in a single study" (Creswell JW 2013).

2.3.1 Rationale

Mixed methods were considered most appropriate for this study due to the complex nature of the data and the study's aim (O'Cathain A 2013;O'Cathain A et al. 2007). Using a combination of quantitative and qualitative methods facilitated 'sense making' defined as "the active process of assigning meaning to ambiguous data". This process enabled a more comprehensive analysis, by providing greater insight into the data than one method could have alone (Bourgeault I et al. 2010;Denzin NK and Lincoln YS 2011;Morse JM 2003). Combining the insights gained from the analyses increased the yield of findings to provide additional nuanced information, that would otherwise have overlooked (O'Cathain A 2013;O'Cathain A et al. 2007).

2.3.2 Design

Cresswell JW and Clark VLP 2007 highlight four aspects to consider when designing a mixed methods study: timing, priority, mixing, and theorising perspectives. The student decided to adopt a sequential explanatory strategy for this study (Creswell JW and Clark VLP 2007). There was no primary data collection in this study which is the main 'timing' consideration, but the data processing did guide qualitative sampling and subsequent analysis, a concept called facilitation (Sandelowski M 1993). The student conducted the quantitative and qualitative analyses sequentially not concurrently (Figure 3).

The priority (or weighting) was given to the quantitative data to provide insights into the more prevalent issues identified in the data, and the qualitative insights supplemented these findings. During the qualitative analyses the student aimed to highlight important, rare, context-specific issues that would be missed by the quantitative analysis alone, to provide an additional perspective on the data. The student compared and linked the findings of the thematic analysis to the quantitative findings.

The qualitative data supported and provided additional insights into the quantitative findings, the quantitative and qualitative data were therefore 'embedded', rather than 'integrated' where the quantitative and qualitative data

are merged, or 'connected', where data are mixed across phases of data collection and analysis (Creswell JW 2013; Creswell JW and Clark VLP 2007). There were no 'a priori' theories informing the design of this research, it was largely inductive in nature

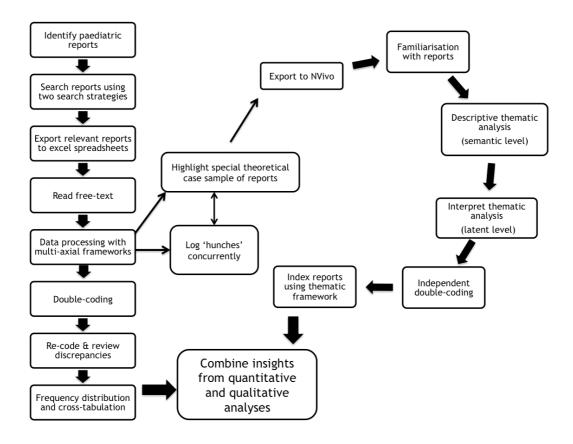


Figure 4 page 40: flow diagram illustrating the processes of mixed methods analysis

2.3.3 Strengthening the analysis

Key attributes of rigorous research include: validity/ credibility, reproducibility, and generalisability (Green J and Thorogood N 2009; Lacey A and Luff D 2001). Triangulating the findings of both quantitative and qualitative analyses increases the internal validity and credibility of results and subsequent confidence in findings (Glik DC et al. 1986). This is in keeping with the positivism and realism perspective: there is a single reality and corroborating and converging multiple

measures can facilitate a more accurate description and understanding of that reality (Bourgeault I et al. 2010).

The reliability of findings, in terms of the consistency and reproducibility of results, was ensured through independent thematic analysis of 100% of the qualitatively sampled reports in addition to double coding of a 20% random sample of reports using the classification frameworks. Kappa statistics were calculated to evaluate inter-rater reliability.

The student and her colleagues (involved in double-coding) undertook root cause analysis training prior to data processing. Weekly team meetings were held to discuss discrepancies between coding, iterations to the classifications, and complex reports. These meetings were recorded and all amendments and decisions were logged providing an audit trail.

Statistical generalisability was not possible with this data, largely as a result of data biases such as under-reporting, although conceptual generalisability, an aim of rigorous qualitative research, was sought (Bourgeault I et al. 2010).

2.4 Ethical Approval

Aneurin Bevan University Health board research risk review committee waived the need for ethical review given the anonymised nature of data (ABHB R and D Ref number: SA/410/13).

Chapter 3: Results

3.1 Vaccination-related incidents

This section presents the results of a mixed methods analysis of vaccination-related safety incident reports involving children in primary care. The student will:

- **3.1.1** provide an overview of the data included and processed, before presenting more in-depth findings from the exploratory quantitative analysis of included data.
- **3.1.2** present the key themes and sub-themes identified from thematic analysis of purposively sampled reports, and describe how those themes relate to each other.
- **3.1.3** combine the insights gained from the quantitative and qualitative analysis to inform a visual model illustrating the weaknesses in the process of routine childhood vaccination, and explain how the qualitative insights relate to the relevant quantitative findings.

3.1.1 Quantitative results

3.1.1.1 Overview of reported vaccination-related incidents in primary care

This section will describe the vaccination-related reports included after data processing (coding with the multi-axial frameworks). Of 2288 reports identified through free-text searches, 1735 reports were included and 553 excluded (see Figure 5). Excluded reports included: those describing non-vaccination related incidents (n=464), system issues which did not result in a patient safety incident (n=65), and those with insufficient free-text information (n=24). The search strategy was designed for maximum sensitivity due to the difficulty of searching these data, hence the low specificity, i.e. the high number of reports, excluded. Kappa statistics of inter-rater reliability were high (k=0.77; p<0.001).

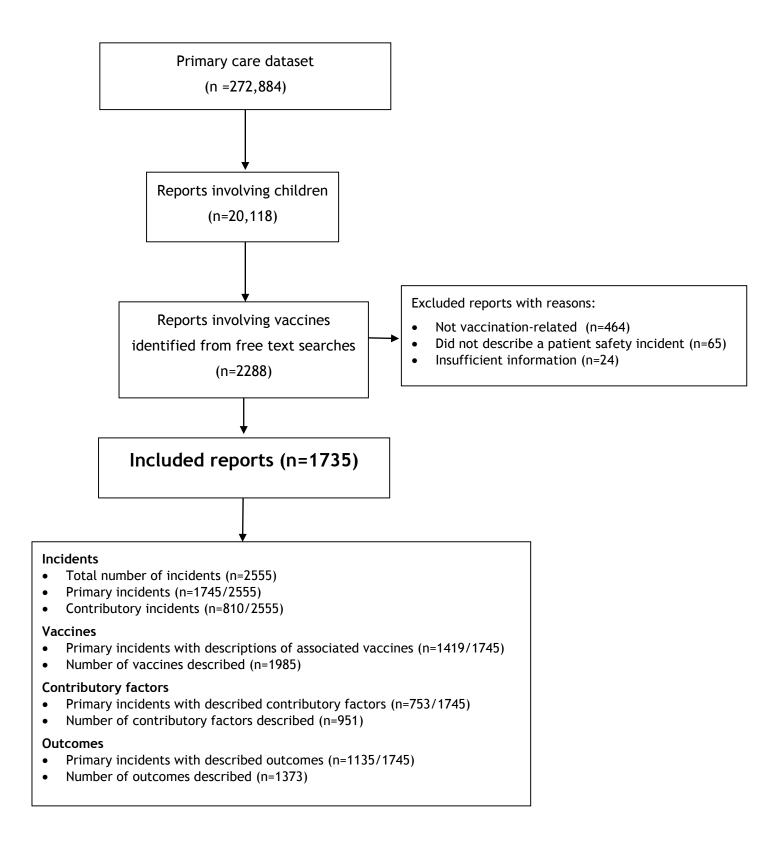


Figure 5 page 44: flow diagram providing an overview of how the included data were retrieved and its content

3.1.1.1.1 Reported ages in vaccination-related incident reports

Primary vaccination-related incidents were most frequently reported in those aged less than one year old, other than a spike of incidents in those aged five years old, the frequency gradually decreased as from those aged one to eight years old, followed by a small increase in incidents in the adolescent years (see Figure 6). This crudely reflects the frequency of vaccinations for each age group specified in the national vaccination schedule.

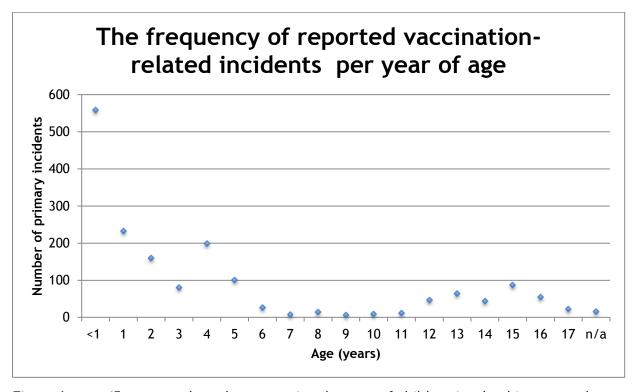


Figure 6 page 45: scatter chart demonstrating the ages of children involved in reported vaccination-related incidents

3.1.1.1.2 Temporal trends in reported vaccination-related safety incidents

The number of reported primary vaccine related incidents increased steeply between the years 2002-2007, they peaked in year 2007 at 279 reports then decreased steadily from 2007 to 2013 but remained above 100-reports a year (see Figure 7). These likely reflect reporting culture rather than the trends in the occurrence of vaccination-related incidents in practice.

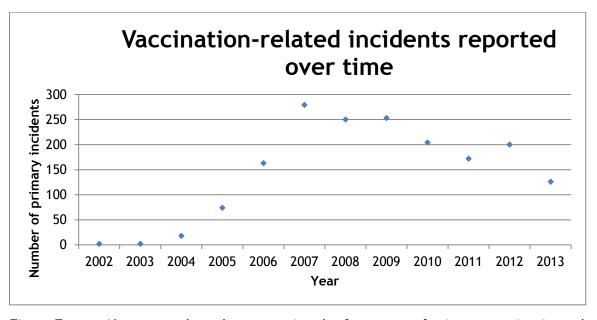


Figure 7 page 46: scatter chart demonstrating the frequency of primary vaccination-related incidents reported over time

3.1.1.3 Vaccines involved in reported safety incidents

The 1745 included primary incidents involved 21 types of vaccine and 1985 vaccines in total (see Table 7 and Figure 8). Most reports of primary incidents (n=1419; 81.3%) described the vaccines involved in the incidents, and some involved multiple vaccines, hence the involvement of 1985 vaccines (Table 7). The most frequently described vaccines were those included in the national immunisation schedule, in particular the MMR¹⁰ (n=361), PCV¹¹ (n=307), DTaP/IPV/Hib¹² (n=241), Men C¹³ (n=226), Hib/ Men C¹⁴ (n=195), DTaP/IPV¹⁵ (n=175), and HPV¹⁶ (n=125) vaccines, account for 82.1% of all vaccines involved in primary incidents (Figure 8).

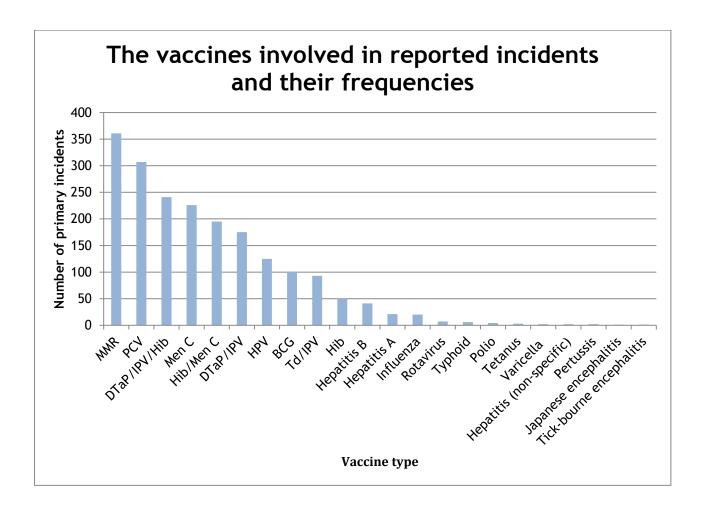


Figure 8 page 47: column chart demonstrating which vaccines were involved in reported primary incidents and their frequency

¹⁰ Measles Mumps Rubella (MMR)

¹¹ Pneumococcal conjugate (PCV)

¹² Diphtheria Tetanus acellular Pertussis/Inactivated Polio/ Haemophilus influenza type B (DTaP/IPV/Hib)

¹³ Meningitis C (Men C)

¹⁴ Haemophilus influenza b/Meningitis C (Hib/Men C)

¹⁵ Diphtheria Tetanus acellular Pertussis/Inactivated Polio (DTaP/IPV)

¹⁶ Human Papilloma Virus (HPV)

Table 7 page 48: the vaccines involved in reported vaccination-related primary incidents

Vaccine	No harm	Low harm	Moderate harm	Death	N codes (%)
Measles Mumps Rubella (MMR)	86	269	6	0	361 (18.2)
Pneumococcal conjugate (PCV)	140	160	4	3	307 (15.5)
Diphtheria Tetanus acellular Pertussis/Inactivated Polio/ Haemophilus influenza type B (DTaP/IPV/Hib)	97	142	1	1	241 (12.1)
Meningitis C (Men C)	103	122	1	0	226 (11.4)
Haemophilus influenza b/Meningitis C (Hib/Men C)	70	123	2	0	195 (9.8)
Diphtheria Tetanus acellular Pertussis/Inactivated Polio (DTaP/IPV)	29	145	1	0	175 (8.8)
Human Papilloma Virus (HPV)	52	59	14	0	125 (6.3)
Bacillus Calmette-Guérin (BCG)	42	52	7	0	101 (5.1)
Tetanus Diphtheria and inactivated Polio (Td/IPV)	15	65	13	0	93 (4.7)
Haemophilus influenza b (Hib)	13	37	0	0	50 (2.5)
Hepatitis B	24	10	7	0	41 (2.1)
Hepatitis A	10	11	0	0	21 (1.1)
Influenza	11	8	1	0	20 (1.0)
Other	6	9	1	0	16 (0.8)
Rotavirus	5	2	0	0	7 (0.4)
Typhoid	2	4	0	0	6 (0.3)
Total	699	1218	57	4	1985 (100)

3.1.1.1.4 Vaccination-related incidents

The 1745 primary incidents were described within 1735 reports since ten reports described more than one independent safety incident. Included reports therefore described 1745 primary incidents, and 2555 incidents in total (when including the 810 contributory incidents) (Figure 5, see Appendix 3.1 for the frequencies of each combination of incidents). Most incidents involved vaccine administration (n=1368). Within this group frequent incidents included administration of the wrong number of doses (n=476), administration of the wrong vaccine (n=318), and administration of vaccines at the wrong time (n=295). Incidents involving vaccine documentation were also frequently described (n=461), in addition to office administration issues (n=216; see Table 8).

Table 8 page 50: the frequency of all incidents (primary and contributory)

Vaccination-related incident	N codes
Vaccination	
Vaccine administration	
Wrong number of doses	476
Wrong vaccine	318
Wrong timing	295
Not administered	83
Wrong dose	57
Out of date	49
Wrong patient	30
Non-specific	23
Contraindicated	21
Wrong site	6
Used needle	5
Wrong storage	4
Wrong route	3
Adverse reaction	148
Reconstitution error	59
Vaccine prescribing and dispensing	33
Insufficient vaccine supply	13
Batch recall	4
Non-specific	1
Documentation	
Records not up to date	261
Record availability	123
Records inaccurate/ unclear	74
Other documentation	2
Administration	
Appointment management	137
Transfer of information	67
Other administration	11
Communication	146
Procedural skills error	88
Referral for vaccination	5
Environmental hazard	5
Medication error	2
Professionalism	2
Treatment decision	1
Lab investigation error	1
Transport errors	1
Insufficient assessment	1
Total	2555

3.1.1.1.5 Factors contributing to vaccination-related incidents

Of the 1745 primary incidents, 753 (43.2%) described 951 contributory factors (see Appendix 3.2 for the frequency of each combination of contributory factors). Staff factors were most frequently described (n=453), these included staff mistakes (n=240) and failure to follow protocols (n=186) (see Table 9). Patient and caregiver factors were also frequently described (n=246). These included their behaviour (n=74), their geographical characteristics (n=64), their knowledge (n=48), and health (n=37). Organisational factors, such as working conditions (n=52), and continuity of care (n=48), education and training (n=36), and organisational protocol failure (n=27), were implicated in 163 incidents. Finally, vaccine factors such as failure (n=36), packaging (n=25), storage (n=25), and design (n=3) contributed to 89 incidents.

Table 9 page 52: the frequency of contributory factors described for each primary incident (NB: *some reports described more than one type of mistake)

Contributory factors - definition	N (%)
Patient/ caregiver factors	
Patient/ caregiver behaviour - the way in which patients or caregivers act or conduct themselves	74 (7.8)
Non-adherence	60
Non-disclosure	12
Other	1
Violence Patient/ caregiver geography the gree where nations live	1
Patient/ caregiver geography - the area where patients live	64 (6.7)
New to area Access difficulties	62 2
Patient health - factors relating to the patient's physical and mental wellbeing	37 (3.9)
Allergy	22
Non-specific	4
Disability	4
Immunocompromised	3
Abnormal coagulation	2
Pregnancy Control of the Control of	2
Patient/ caregiver knowledge - insufficient knowledge or inadequate application of knowledge	48 (5)
Looked-after child - children not in the care of their parents e.g. in foster care	18 (1.9)
Patient/ caregiver language - patient or caregiver unable to communicate in English	5 (0.5)
Staff factors	
Mistake - cognitive lapses	*240 (25.2)
Non-specific mistake	139
Similar vaccine appearances	45
Distraction	22
Misreading	18
Inattention	10
Similar patient names	9
Failure to follow protocol - not adhering to organisational guidelines	186 (19.6)
Knowledge - insufficient knowledge or inadequate application of knowledge	19 (2)
Fatigue/ stress - extreme tiredness, mental or emotional strain	5 (0.5)
Other factors	3 (0.3)
Equipment/ vaccine factors	
Failure of equipment/ vaccine - the equipment or vaccine is faulty	36 (3.8)
Equipment/vaccine packaging - the packaging is impractical inadequate or faulty	25 (2.6)
Equipment/vaccine storage - inadequate impractical storage	25 (2.6)
Poor equipment/vaccine design - the design is impractical, inadequate or faulty	3 (0.3)
Organisational factors	
Working Conditions - factors relating to the work environment	52 (5.5)
Continuity of care - issues with the co-ordination of services	48 (5)
Education and training - insufficient education and training of staff	36 (3.8)
Inadequate guidelines or protocols - existing guidelines not fit for purpose	27 (2.8)
Total	951 (100)

3.1.1.1.6 Severity of harm resulting from vaccination-related incidents

Most reports (n=1077; 61.7%) described harm of some sort including: three deaths, 67 incidents of moderate harm, and 1007 incidents of low harm (Figure 9). However the pre-allocated harm severity entered by reporters differs considerably to the harm severity using the WHO definitions (Table 10). Prior to classification using the WHO definitions, included primary incidents were mostly classified (by reporters) as not harmful (n=1390; 79.7%). Many (n=775) primary incidents were upgraded, 739 were upgraded from no harm to low harm, 35 from no harm to moderate harm, and 1 incident from moderate harm to death. Fewer were downgraded in terms of harm: 27 were downgraded to no harm and 21 to low harm. The differences are as a result of numerous primary incidents being up-graded and downgraded in terms of harm severity-based on the outcomes described by reports.

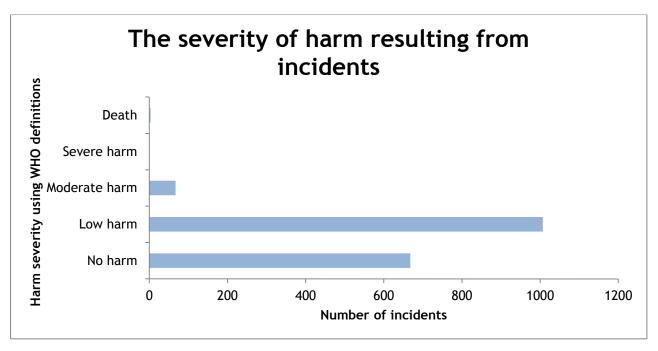


Figure 9 page 54: bar chart demonstrating the severity of harm resulting from incidents

Table 10 page 54: demonstrates how many primary incidents had their allocated harm severity upgraded (in blue) and downgraded (in green), and how many kept their preallocated harm severity (yellow) (*1 report of moderate harm was upgraded to death)

Re-coded harm		Reporter-allocated severity in reports								
severity using WHO			Moderate	Severe						
definitions	No harm	Low harm	harm	harm	Death	N codes				
No harm	641	24	2	1	-	668				
Low harm	739	247	21	-	-	1007				
Moderate harm	10	25	32	-	-	67				
Severe harm	-	-	-	-	-	0				
Death	-	-	1*	-	2	3				
N codes	1390	296	56	1	2	1745				

3.1.1.7 Outcomes of vaccination-related incidents

Most primary incidents (n=1135; 65%) described outcomes and many described multiple outcomes; therefore 1373 outcomes are described for 1135 primary incidents. The most frequently described outcomes were patient inconvenience $(n=801)^{17}$ such as: receiving unnecessary treatment (n=481) and requiring additional treatment (n=379). Other outcomes described included clinical patient harm (n=205) such as injuries (n=72) and fainting (n=64), and exposing the patient to risk (n=139), for example by leaving them vulnerable to vaccine preventable diseases (n=108).

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¹⁷ Note some reports described multiple types of patient inconvenience

3.1.1.2 Primary Incident Types

This section will present a more in-depth analysis of the incident types that were harmful and described most frequently. The primary incident types described by reports mostly involved vaccine administration incidents (n=1282/1745; 73.5%), however adverse vaccine reactions (n=146), procedural issues (n=57), and communication incidents with patients and caregivers (n=51) were also evident. For the purposes of this study adverse drug reactions included any unintended, undesirable, or unexpected effects of prescribed medications or of medication errors (Joint Commission Resources 2005). The relationship between these incident types, time of report submission, harm severity, contributory factors and the vaccines involved are presented in Tables 11-14. Appendices 3.1 and 3.2 demonstrate the frequencies of combinations of incidents and contributory factors.

3.1.1.2.1 Vaccine administration

Vaccine administration primary incidents included administering the wrong number of doses (n=476), administering vaccines at the wrong time (n=294), and administering the wrong vaccine (n=249). These account for 79.5% of vaccine administration primary incidents and following the Pareto Principle (that 80% of the problem can be addressed by focusing on 20% of the issues - which was originally a principle of welfare economics) this section will focus on these three types of administration primary incidents (NHS Scotland 2015).

Table 11 page 57: the number primary incidents reported each year within each incident type

	Number of reports per incident type per year													
Primary incident types	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	N/A	N codes
Vaccination														
Administering														
Wrong number of doses	1	2	1	23	42	100	60	75	51	38	53	29	1	476
Wrong timing	-	-	2	8	24	47	52	35	38	28	34	26	-	294
Wrong vaccine	-	-	12	15	28	45	34	29	26	21	19	19	1	249
Not administered	-	-	-	2	8	4	35	9	5	6	7	4	-	80
Wrong dose	-	-	=	6	7	7	4	8	9	6	7	3	-	57
Out of date	-	-	-	2	3	7	6	12	8	5	4	2	-	49
Non-specific	-	1	-	-	6	7	3	-	1	1	4	-	-	23
Contraindicated vaccine	-	-	-	-	2	1	3	5	-	-	8	1	-	20
Wrong patient	-	-	2	3	1	-	=	1	2	3	3	1	-	16
Wrong site	-	-	-	-	-	-	1	-	4	-	1	-	-	6
Used needle	-	-	-	-	-	-	-	2	-	1	1	1		5
Wrong storage	-	-	-	-	-	1	2	1	-	-	-	-	-	4
Wrong route	-	-	-	-	-	-	-	1	-	-	1	1	-	3
Adverse reaction	-	-	1	2	10	12	8	26	31	19	29	8	-	146
Vaccine prescribing and dispensing	-	-	=	-	2	11	3	2	1	4	1	2	-	26
Reconstitution error	-	-	=	=	-	-	3	=	-	1	=	-	-	4
Batch recall	-	-	=	=	-	-	=	4	-	-	=	-	-	4
Insufficient supply	-	-	-	-	-	-	1	-	-	-	-	-	-	1
Non-specific	-	-	-	=	-	1	-	-	-	-	-	-	-	1
Documentation														
Records not up to date	-	-	-	2	3	12	7	4	8	20	5	5	-	66
Records inaccurate/ unclear	-	-	-	1	4	4	3	4	3	4	5	3	-	31
Records unavailable	-	-	=	=	-	-	2	-	2	-	-	2	-	6
Procedural error	-	-	-	4	9	6	10	11	5	5	4	3	-	57
Administration														
Appointment	-	-	-	-	4	2	5	9	5	1	6	7	-	39
management Transfer of	-	-	-	-	2	2	1	4	2	-	-	4	-	15
information Other	-	-	-	-	-	1	-	-	-	-	-	1	-	2
Communication														
With patients/	-	-	-	6	7	8	7	8	-	6	5	4	-	51
caregivers Between healthcare	-	-	-	-	-	-	-	1	-	1	-	-	-	2
professionals Environmental hazard	_	-	-	-	1	_	_	2	1	1	-	-	-	5
Referral for	-	-	-	-	-	_	-	-	1	_	2	-	-	3
vaccination Professionalism	-	-	-	-	-	-	-	-	1	-	1	-	-	2
Medication error	-	-	-	-	-	-	-	-	-	1	-	-	-	1
Transport	-	-	-	_	-	1	-	-	-	_	-	-	-	1
•			18								200	126		

Table 12 page 58: the frequency and severity of harm described for each age group and for each primary incident type

	Severity of harm				
	No	Low	Moderate	Death	N codes
Age	harm	harm	harm	Death.	1, codes
Under 28 days	15	7	3	-	25
1 month to 1 year	307	246	13	3	569
2 to 4 years	206	433	9	-	648
5 to 11 years	41	133	5	-	179
12 to 17 years	99	188	37	-	324
Incident Type					
Vaccination					
Administering	438	824	17	3	1282 (73.5)
Wrong number of doses	28	447	1	-	476
Wrong timing	239	44	8	3	294
Wrong vaccine	97	150	2	-	249
Not administered	14	65	1	-	80
Wrong dose	26	31	-	-	57
Expired vaccine	3	46	-	-	49
Non-specific	10	13	-	-	23
Contraindicated vaccine	14	5	1	-	20
Wrong patient	2	14	_	-	16
Wrong site	1	2	3	-	6
Used needle	3	2	-	-	5
Wrong storage	-	4	-	-	4
Wrong route	1	1	1	-	3
Adverse reaction	-	103	43	-	146 (8.4)
Prescribing and dispensing	26	-	-	-	26 (1.5)
Reconstitution error	-	4	-	-	4 (0.2)
Batch recall	4	-	-	-	4 (0.2)
Insufficient supply	1	-	-	-	1 (0.1)
Non-specific	1	-	-	-	1 (0.1)
Documentation	97	5	1	-	103 (5.9)
Records not up to date	62	3	1	-	66
Records	29	2	-	-	31
inaccurate/unclear					
Record availability	6	-	-	-	6
Administration	50	4	2	-	56 (3.2)
Appointment management	36	3	-	-	39
Transfer of information	12	1	2	-	15
Other	2	-	-	-	2
Procedural errors	4	53	-	-	57 (3.3)
Communication	44	9	-	-	53 (3)
With patients or caregivers	42	9	-	-	51
Between HCPs	2	-	-	-	2
Other	3	5	4	-	12 (0.7)
N codes	668	1007	67	3	1745

Table 13 page 59: the frequency of contributory factors described for the key primary incident types

	Primary incident type										
Contributory factors	Wrong number of doses	Wrong vaccine	Wrong timing	Adverse reaction	Procedural error	Communication with patients and caregivers					
Patient/ caregiver factors											
Patient/ caregiver behaviour											
Non-adherence	1	-	1	2	39	-					
Non-disclosure	5	-	1	2	-	-					
Other	-	-	1	-	-	-					
Patient/ caregiver geography											
New to area	36	1	13	-	-	-					
Access difficulties	-	-	2	-	-	-					
Patient health											
Allergy	-	-	-	18	-	-					
Other health issues	-	-	1	2	-	-					
Disability	-	-	1	-	-	-					
Coagulation problems	-	-	-	1	-	-					
Pregnancy	-	1	-	-	-	-					
Patient/ caregiver knowledge	40	-	4	-	-	1					
Looked-after child	8	1	3	-	-	1					
Patient/caregiver language	1	-	4	-	-	-					
Staff factors											
Mistake											
Non-specific mistake	47	35	19	-	-	4					
Similar vaccine names	1	37	-	-	-	-					
Misread	10	3	1	-	-	-					
Distracted	2	8	2	-	-	1					
Inattention	3	4	-	-	-	-					
Similar patient names	4	-	-	-	-	-					
Failure to follow protocol	63	37	35	1	-	4					
Knowledge	3	6	3	-	1	-					
Fatigue/stress	-	5	-	-	-	-					
Other factors	1	-	1	-	-	-					
Equipment/ vaccine factors											
Failure of equipment/ vaccine	-	1	2	-	10	-					
Equipment/ vaccine storage	-	10	-	-	-	-					
Equipment/ vaccine packaging	1	3	-	-	-	-					
Organisational factors											
Working conditions	10	14	12	-	-	-					
Continuity of care	15	-	22	-	-	-					
Education and training	6	10	11	-	1	1					
Inadequate protocol or guidelines	3	3	3	-	-	1					
N codes	260	179	142	26	51	13					

Table 14 page 60: the vaccines involved in the most frequently reported primary incident types

Type of administration		Vaccine type													
incident	MMR	PCV	DTaP/IPV/Hib	Men C	Hib/Men C	DTaP/IPV	HPV	BCG	Td/IPV	Hib	Hep B	Other	Hep A	Flu	N codes
Wrong number of doses	156	75	63	49	77	87	8	20	36	16	0	5	3	1	596
Wrong timing	48	91	83	37	29	7	16	14	2	2	21	6	1	0	357
Wrong vaccine	41	85	56	87	61	49	8	3	23	1	1	5	6	4	430
Not administered	12	2	5	13	4	3	1	6	1	28	1	0	1	0	77
Wrong dose	12	1	1	7	3	1	2	9	0	0	5	1	5	3	50
Expired vaccine	12	0	1	2	4	7	1	5	1	3	1	1	2	2	42
Contraindicated vaccine	2	1	0	1	0	2	0	5	1	0	0	2	0	2	16
Wrong patient	7	1	4	2	1	1	2	0	1	0	0	0	0	0	19
Other	2	5	2	3	4	1	2	5	0	0	2	3	0	2	31
Adverse reaction	19	14	9	5	2	9	52	4	20	-	-	2	-	-	136
Communication with patients or caregivers	25	4	1	-	-	1	10	2	1	-	-	1	-	1	46
Procedural error	8	3	2	3	3	3	3	3	1	-	-	-	-	1	30

3.1.1.2.2 Wrong number of doses administered

Children receiving the wrong number of doses were mostly aged less than 6 years old. These incidents were frequently reported in those aged 0-2 years (n=141), decreasing in those aged three to four years (n=23), and increasing again in those aged four to six years (n=131). (Figure 10)

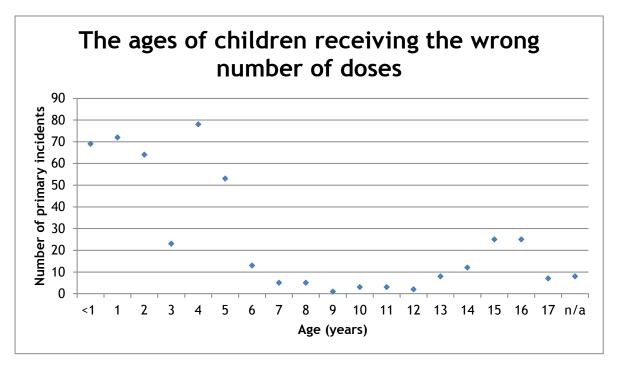


Figure 10 page 61: a scatter chart illustrating the ages of children receiving the wrong number of doses

Most incidents of administering the wrong number of doses were harmful (n=448; 94.1%); this was typically because the child had received an unnecessary additional vaccination and thus met the criteria for 'low harm' (Table 12). However, one of these incidents, where a child suffered an adverse reaction and required hospitalisation, resulted in moderate harm. The vaccines often involved in these incidents were MMR (n=156), DTaP/IPV (n=87), and Hib/Men C (n=77), which are given routinely and in multiple times as part of the childhood vaccination schedule (Table 14).

These incidents were often preceded by other contributory incidents (see Appendix 3.1 for the frequencies of each combination of incidents). Documentation failures such as out of date (n=128), unavailable (n=45,) or inaccurate documents (n=15) were frequently implicated. Three forms of vaccination documentation were described in reports: parental held records (red books), GP records, and child health records (in a public health

repository). Discrepancies between these records accounts for some of the various documentation failures described above i.e. not all three records were available, accurate, or checked prior to vaccine administration.

Multiple vaccine administration incidents were occasionally described in the same report (see Appendix 3.1 for the frequencies of incident combinations). For example, 50 children received the wrong number of vaccine doses, because the wrong vaccine had been administered (a contributory incident). Similarly 6 children received the wrong number of doses because they received someone else's vaccination i.e. the wrong child was vaccinated. Other contributory incidents included: communication errors (n=26), and issues with appointments (n=19).

Patient and caregiver factors frequently contributed to these incidents (Table 13, see Appendix 3.2 for the frequency of each combination of contributory factors). For example, patient and caregiver knowledge-such as not knowing which vaccines were needed or had been previously received-was implicated in 40 of these incidents. Other patient and caregiver factors included being new to the area and general practice (n=36), and eight incidents were partly the result of a child being 'looked after' (in Local Authority care). Staff factors contributing to incidents include: not following protocols (n=60) (intentionally or unintentionally) such as failing to check all the appropriate documentation prior to administration; and mistakes (n=67) such as misreading vaccine names (n=10). Fewer organisational issues were described, however poor continuity of care (n=15) and insufficient staffing (n=10) were described as contributing to some incidents.

Example reports of administration of the wrong number of vaccine doses

Example 1: "Patient presented with stepmother for pre - school booster. Written consent from father was brought but parental held record was not available. Nurse explained she was giving repevax [DTaP/IPV] and MMR. The following day stepmother called expressing concern that MMR had already been given in 2004. Incomplete documentation of initial dose of MMR."

Example 2: "Child placed with adoptive parents who were advised by Social Worker to attend the GP to complete primary vaccinations. Attended surgery with parental held records but no family practice records were available. Only two immunisations were recorded in the parental held record. Immunisation given with consent. Later informed by Social Services that child had already completed her primary immunisations. Family practice records checked and confirmed above."

Example 3: "Patient received the third primary immunisation twice, in error, once in Ghana and once at the health centre. Mother failed to notify the Health Visitor of the first immunisation."

3.1.1.2.3 Wrong timing

Most children receiving vaccines at the wrong time were aged less than 1 year old (n=175). The frequency of these incidents decreased steeply to 43 and 27 in those aged 1-2 years and 2-3 years respectively. The number of incidents then plateaued as the children increased in age (Figure 11). Administration of vaccines at the wrong time included incidents where vaccination had been delayed, and incidents where the vaccine has been administered contrary to the national recommended schedule for example, administering vaccines within two weeks of each other rather than the recommended four.

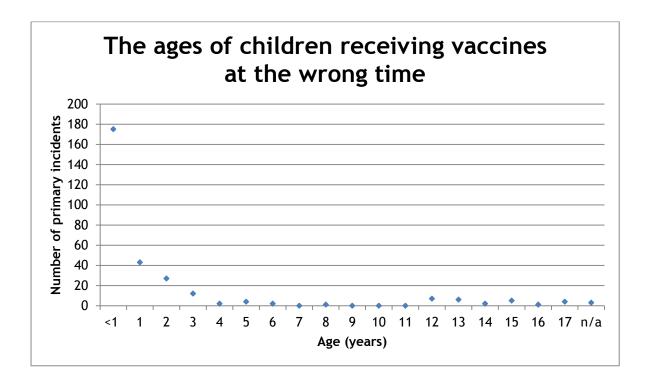


Figure 11 page 64: scatter chart illustrating the frequency of vaccine-timing incidents in each age group

Vaccines typically involved were Pneumococcal conjugate (PCV) (n=91, 31%), Diphtheria Tetanus acellular Pertussis/Inactivated Polio/ Haemophilus influenza type B (DTaP/IPV/Hib) (n=83, 28%), and the MMR (n=48, 16%) (Table 14).

The proportion of incidents resulting in harm was less for this group (wrong timing) than other vaccine administration incidents; however incidents in this group were more harmful in terms of severity (Table 12). Of 294 incidents, 55 (18.7%) were harmful: eight of these were cases of moderate harm, and three were deaths. The children suffering moderate

harm tended to be newborns who required prophylactic Hepatitis B or BCG¹⁸ vaccination but in whom administration was delayed or not in line with current guidelines- therefore exposing these infants to unnecessary risk. The three children who died had pneumonia and meningitis, and were all delayed in receiving the appropriate vaccinations, which if received may have prevented infection. The 44 incidents of low harm typically described children who required additional vaccinations for adequate immunity.

Incidents contributing to these timing issues involved: appointment management difficulties (n=60), documentation failures (n=31), communication errors (n=21), transfer of documentation between care settings (n=16), and administering the wrong vaccine (n=12) (see Appendix 3.1 for the frequencies of each combination of incidents). Poor continuity of care (n=22) -at organisational level- such as health visitors not receiving birth notifications from secondary care was described as contributing to these incidents (Table 13, see Appendix 3.2 for the frequency of each combination of contributory factors). These incidents tended to occur in patients who were new to the area, looked-after, who had (or whose caregivers had) poor knowledge. Staff deviating from protocols (n=32) and making mistakes (n=22) were also implicated.

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¹⁸ Bacillus Calmette-Guérin (BCG)

Examples of reports describing vaccination timing-related incidents

Example 4: "An infant died from a streptococcal pneumonia - which could have been prevented if the child had received childhood immunisations. The mother stated she was not aware that her child should be immunised and the child was not registered at a family practice until *** Identified areas of concern include: - the management of the child immunisation processes, family practice registration processes and notifying child health of non-registered patients."

Example 5: "Patient's relative contacted health visitors regarding her child's immunisations, she reported she had not received any appointments for her child's third primary immunisations. Child health computer had recorded wrongly that the child had his third immunisations on the same day as he had his second immunisations. The patient received his third primary immunisations late because of this. Child health would not have been aware of this if the parent had not contacted the service."

Example 6: "Patient was scheduled for Hib/MENC vaccine; staff checked his immunisation record and became aware that he already had this immunisation. At this point I made an error. I told the mother that we could give the MMR / Prevenar. Mother's English is not perfect and she agreed. As I came to record the immunisation, I realised my error there was a two-week gap between immunisations not 4."

3.1.1.2.4 Wrong vaccine administered

Children reported as receiving the wrong vaccine were largely aged less than one year old (See Figure 12). Many incidents (n=249) involved administration of the wrong vaccine, most of these were harmful (n=152; 61%) as children who received the wrong vaccine had often received an unnecessary treatment and then also required additional treatment (the vaccine that was originally required) (see Table 12).

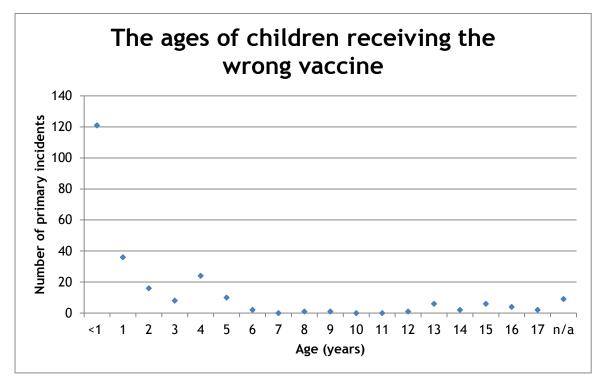


Figure 12 page 67: scatter chart demonstrating the frequency and ages of children receiving the 'wrong vaccine'

The vaccines most frequently involved in these incidents include: Men C (n=87), PCV (n=85), and Hib/ Men C (n=61) (Table 14). Often when a single Men C vaccine was scheduled e.g. at 3 months children wrongly received either PCV (n=40) (scheduled at 2 and 4 months), or a Hib/ Men C combination vaccine (n=31).

These incidents were not often preceded by other incidents such as documentation failures (n=18), and vaccinating the wrong patient (n=6) (see Appendix 3.1 for the frequencies of each combination of incidents). However incidents were often the result of staff failures such as: failing to follow protocols, and mistakes (n=87) such as confusing similar vaccines (n=37). Ambiguous packaging (n=13) was also described as contributing to

these mix-ups (Table 13, see Appendix 3.2 for the frequency of each combination of contributory factors).

Example reports of administration of the wrong vaccine

Example 7: "Mother took her five-month-old baby to her GP for his second DTaP/IPV/Hib vaccination. The Staff Nurse administered the wrong injection because she did not consult his medical records. The baby was given an MMR vaccination that should not be given until he is 13 months old. Staff Nurse says she was distracted during the appointment. The Nurse will re - train and demonstrate her competency through supervision."

Example 8: "The patient attended for booster immunisations. I proceeded to give the vaccination and documented the batch number. Later I realised that I had given the patient the wrong immunisations. I had confused the patient with another patients 'who was also due unscheduled immunisations."

3.1.1.2.5 Adverse reactions to vaccines

Adverse reactions were described as the primary incident by 146 reports: all of these were harmful including 103 cases of low harm and 43 cases of moderate harm (Table 12). Most of these were in adolescents (Figure 13). The children suffering moderate harm were sent to the emergency department with various symptoms: left sided weakness, altered consciousness, rash, vomiting, shortness of breath, slurred speech, dilated pupils, fevers, and seizures. Some needed adrenaline and some had head injuries as a result of losing consciousness. Most of the moderately harmful reactions (n=26/43) were the result of either HPV (n=13) or Td/IPV¹⁹ (n=13) vaccines.

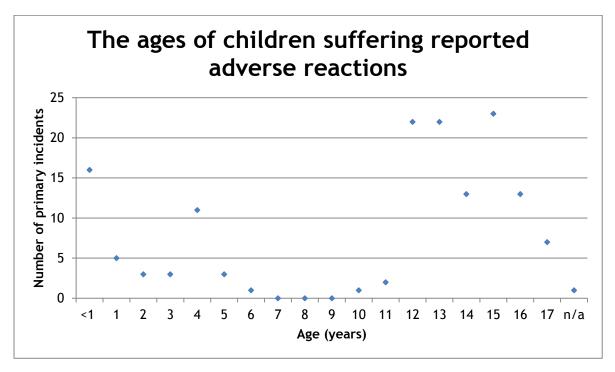


Figure 13 page 69: scatter chart illustrating the frequency and ages of children reported as suffering adverse reactions to vaccines

Most reports (n=117, 80%) described the vaccines involved in each incident, of which there were 136 (Table 3.8). Approximately a third of adverse reactions described were from HPV vaccination (n=50) in adolescents. Other frequently described vaccines included: Td/IPV (n=20), MMR (n=19), and PCV (n=14). Some children had pre-existing medical conditions that were not communicated to the health care professionals either by the child or the caregiver prior to vaccination (particularly in schools where caregivers typically provide this information on consent forms sent via their children).

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¹⁹ Tetanus Diphtheria and inactivated Polio (Td/IPV)

Communication incidents contributed to some (n=4) of these incidents (see Appendix 3.1 for the frequencies of each combination of incidents).

Patient allergy was the most frequently described contributory factor (n=18). This includes patients with known allergies and those in whom the allergy was unknown prior to the incident (Table 3.7, see Appendix 3.2 for the frequency of each combination of contributory factors). Other patient contributory factors included having medical conditions (n=3) that predispose to adverse vaccine reactions e.g. immune thrombocytopenia.

Example reports of adverse reactions

Example 9: "Administered hepatitis vaccine 1st dose at 11:15am. At 11:30 he complained of feeling extremely unwell, faint and nauseous. His blood pressure was 67 / 45. He began to shake uncontrollably complaining of feeling hot then very cold. He then complained of feeling tired and wanting to sleep and had a severe headache; he kept slipping in and out of consciousness but came to when I shouted his name. I asked school staff to call for an ambulance and to contact his parents. His mum arrived and informed me he had allergies to cow milk, tomatoes, citrus fruits, however had not put this information on the vaccine consent form, although these allergies were not contraindications. Mum also stated C recently attended A&E with similar symptoms. Taken to A&E by ambulance. Yellow Card completed."

Example 10: "Child received revaxis [Diphtheria Tetanus acellular pertussis and Inactivated Polio] immunisation at school. Has known nut allergy but never been allergy tested and has no Epi pen. 45 minutes later he returned with slurred speech, disorientated and feeling dizzy. He appeared very pale. Assisted to lay down with legs raised and began to complain of a strange tight feeling in his throat and his lips appeared to be blue in colour."

Example 11: "Telephone call received today by parent expressing concern that her daughter had reacted to her third HPV vaccination. Mum concerned that she felt that she had not received enough information prior to the injection. Mum reported her symptoms as - double vision with peripheral vision loss, nausea, fatigue and headache."

3.1.1.2.6 Vaccination-related procedural errors

There were 57 procedural errors e.g. scratching a child with the vaccine needle, and 50 of these were in children aged less than five years old. Most (n=53, 93%) of these were harmful as they resulted in injuries (n=42) or a child requiring an additional vaccination or treatment (n=11) (Table 3.6). No contributory incidents were described; however, most reports (50, 88%) described contributory factors, of which there were 51 (Table 3.7, see Appendix 3.2 for the frequency of each combination of contributory factors). Most incidents were the result of poor patient or caregiver adherence (n=39) e.g. not holding the child appropriately during the procedure (hence most of these incidents occurring in those aged less than five years old), some were the result of equipment issues (n=10), and one was the result of poor staff knowledge (n=1) and education (n=1).

Example reports of procedural errors

Example 12: "Patient attended for MMR father relaxed his hold of patient during administration of immunisations causing a scratch to his right arm and MMR to be expelled from syringe doctor contacted further dose of MMR given with parents' agreement."

Example 13: "Whilst giving MMR vaccination the child pulled the syringe out I administered the vaccination as per policy but the child had two puncture sites on his thigh parents witnessed incident and are aware."

Example 14: "Patient attended surgery together with mother for her PC vaccine mother was asked and also shown to hold patients arms and legs on giving the injection patient grabbed my right hand causing the needle to scratch her near the injection site scratch was cleaned and a plaster applied."

Example 15: "When attempting to give patient his meningitis C vaccine, his mother who was crying loosened her grab of him. The baby wriggled and unfortunately sustained a linear scratch about an inch long. Mum became even more upset. I reported the incident to the Health visitor who had a chat with mother and tried to calm her down. Mother eventually relaxed and agreed for me to proceed with vaccination baby."

3.1.1.2.7 Communication with patients/ caregivers about vaccination

Communication issues with caregivers and patients-such as failure to gain adequate consent for vaccination-were described as the primary incident type in 51 reports and 39 of these were related to consent. Most incidents involved children aged less than five years old (n=28, 55%) and were not harmful (n=42, 82%) (Table 3.6). Low harm was described by nine reports largely due to parental distress after their child received a vaccine they did not wish their child to receive.

Most reports described the vaccines involved (n=44, 86%), and almost half of these communication incidents were about the MMR vaccine (n=25) (Table 3.8). The HPV vaccine was also involved in some (n=10) communication incidents.

Example reports of communication incidents

Example 16: "Patient was brought by her mother for pre - school booster injections for pertussis, diphtheria, tetanus and polio. She was also given an injection for measles, mumps and rubella which was not on the appointment card and mother had not intended she should have. Immunisation was given by 2 practice nurses. Mother held child during procedure. The mother of the child had contacted the police when she realised what had happened, claiming her child had been assaulted as a result. At that time the police had taken advice from the CPS and as a result they would not be pursuing charges. Subsequent the member of staff has received a summons to appear before the local magistrate's court. It appears the mother is pursuing this matter privately."

Example 17: "During HPV vaccination sessions, clients consent form was signed by her mother indication that she did not want her vaccination. Her form had not been separated from positive consent forms. I then signed form indication client has refused."

3.1.2 Thematic analysis of vaccination-related reports

Three overarching themes were identified from reports: vulnerability to vaccination incidents; responsibility for childhood vaccinations; and fragmented or substandard services. These will be presented, along with their respective sub-themes, and their overlap, and illustrated in a visual model (Figure 14). The association between these themes and quantitative findings will be discussed further in the mixed methods section. To support the findings discussed in this section, edited extracts of free-text have been provided. These examples include the more extreme and the more typical cases to illustrate the breadth of each theme.

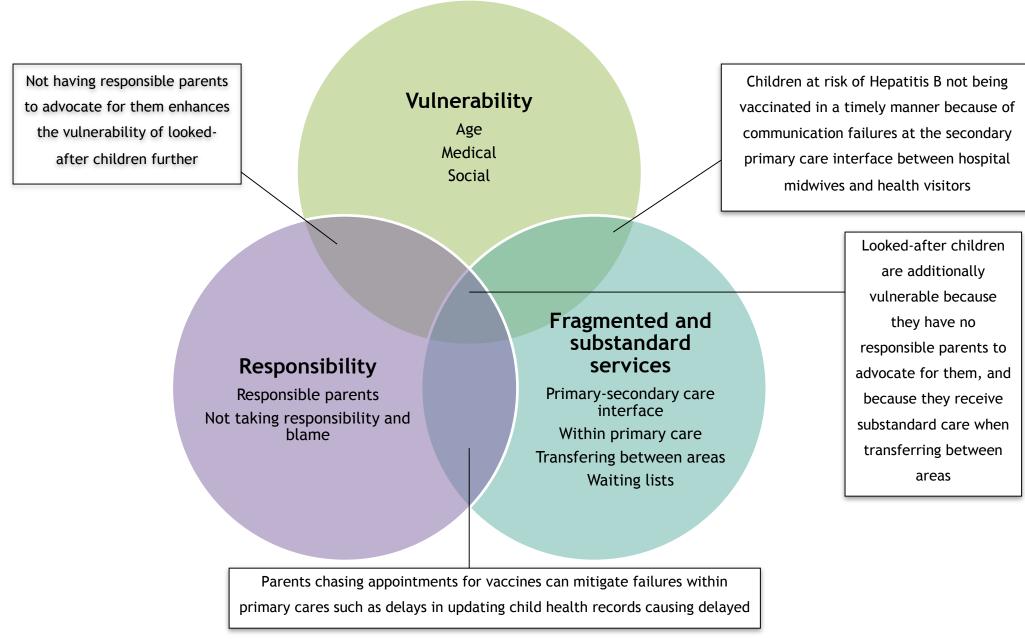


Figure 14 page 74: illustrating the themes identified, their respective sub-themes and how they overlap

3.1.2.1 Vulnerability

3.1.2.1.1 Age-related vulnerabilities

Age featured in various forms: children received adult vaccine doses, or vaccines contraindicated in their age group; additionally there were issues around consent and acknowledging Gillick competence i.e. the child's capacity to provide consent.

Examples

Example 18: "Patient reported that in her opinion her right to choose whether to be immunised had been ignored by a staff member (despite her parent consenting to the vaccine), and that she had been restrained physically by 1 staff whilst another staff approached from behind and administered without her consent."

Example 19: "Child patient was administered with Adult Hepatitis A vaccine (Havrix Monodose) instead of the paediatric dose (Havrix Junior Monodose)."

Example 20: "During the morning I gave 50 vaccinations to adults at approximately 10:30 am - a child *** came for vaccination I had assumed that he was eligible for the older child dose of 0.5mls Pandemrix [Swine flu] - and I gave this. This was an error - since at 9 and under he should have had 0.25mls Pandemrix."

Example 21: "4 year old attended with parents for seasonal flu vaccine. Prescription issued within GP surgery and collected from pharmacy. On reading patient information states that the vaccine (Enriza) is not to be administered to children under 5 due to increase in convulsions."

3.1.1.1.2 Medically vulnerable children

Medically vulnerable children were repeatedly described as experiencing vaccination-related incidents, including children who were pregnant, receiving immunosuppressant treatment for cancer, or at risk of Tuberculosis, HIV^{20} or Hepatitis B.

Pregnant adolescents were receiving vaccines that are contraindicated in pregnancy because the young adult did not disclose the pregnancy, they are unaware of the pregnancy themselves, or the healthcare professional forgot that pregnancy was a possibility in this age group (an area of overlap with the age subtheme). Such incidents did not feature frequently but are important to consider due to the high incidence of teenage pregnancies and the potential for harm to occur to mother and baby.

Examples

Example 22; "Pupil was vaccinated twice when pregnant. Asked at both sessions if possibility of pregnancy and not declared at the time of vaccination. Pupil did not know she was pregnant."

Example 23: "Patient was booked in for a whooping cough vaccine, patient attended appointment with practice nurse, practice nurse administered a HPV vaccine in error, and patient was 28 weeks pregnant. . Patient wished to make a formal complaint, full investigation in progress."

The potential harm to immunosuppressed children receiving contraindicated vaccines is also significant. These incidents were partially a reflection of the inadequate knowledge of healthcare professionals and failures at the specialist secondary- primary care interface (this will be discussed further within the fragmented and substandard services section 3.1.2.3). Caregivers (with first-hand knowledge of their child's medical history were described both as trying to prevent these incidents (raising concerns) and contributing to them by disclosing incorrect information to healthcare professionals (this responsibility theme will be discussed further in the next section).

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²⁰ Human Immunodeficiency Virus (HIV)

Example

Example 24: "Baby received routine immunisations in baby clinic when consultant baby is under had written to surgery requesting baby not receive immunisations until he is one year post chemotherapy. Mother believed baby should have immunisations and he had received two flu vaccines between two sessions of chemotherapy. GP had seen letter from consultant and had put a note on baby record. Note on baby records was not dated and was believed to be automatic warning."

Children who were at risk of Tuberculosis and Hepatitis and who therefore required vaccination were often involved in incidents. These vaccines are not routinely scheduled vaccinations, the need for them is typically identified antenatally, they are arranged by the hospital postnatally and this requirement is then communicated to community midwives or health visitors. Looked-after children requiring these vaccines appeared even more vulnerable to incidents; such children also feature in the subsequent sub-theme of social vulnerability. A subset of these children at-risk of Tuberculosis and Hepatitis B had HIV positive mothers; the BCG vaccine is contraindicated in these children until they have been confirmed as HIV negative - these additionally vulnerable new-borns were occasionally described as 'slipping through' the safety net and almost all receiving BCG vaccines. Poor care co-ordination between secondary care and health visiting, or midwifery and health visiting, were repeatedly described as the reason for these 'near-misses' (an area of overlap with the fragmented and substandard services section 3.1.2.3).

Examples

Example 25: "A mother brought her baby to the BCG clinic at ****. During the consultation I was made aware by the mother that she was HIV positive. This is a contraindication to the baby receiving BCG vaccination until baby has been screened and given the all clear by the paediatricians. This was explained to the mother and reasons given. I gave apologies for the inconvenience and advised her that I would be looking into the reasons for her receiving the appointment. Health visitor did not adhere to agree process by not indicating on the form that the mother was HIV positive."

Example 26: "At Lac Review on 14.4.11 was highlighted that child overdue 2nd Hep B vaccine no entry made in red book, discharge note did not indicate Hep B given at birth, no indication GP made aware of follow up. There appears to have been no contact with the GP surgery to indicate this would need following up at all."

Example 27: "Baby discharged with request to give a BCG vaccination. Appointment arranged and consent obtained from the mother using an interpreter. BCG given but when the records were updated it was noted that the ward had given the BCG prior to discharge."

Example 28: "Audit of outcome for baby born to Hepatitis B positive mother received in child health department 6 / 7. Midwife written on bottom 'sincere apologies for not completing paperwork at delivery'. Notification received so late after delivery, very real risk that child would not have received 2nd and 3rd dose."

3.1.1.1.3 Socially vulnerable children

A proportion of vulnerable children such as looked-after children, asylum seekers, and immigrants suffered vaccination-related incidents as a result of these vulnerabilities. The systems in place failed these vulnerable children who often had inaccurate or no medical records, had difficulty accessing care, and some had difficulty communicating with healthcare professionals in English. For example one looked-after child had no vaccine records because whilst in her mother's care the vaccines had been administered at a private health centre. Other looked-after children suffered incidents because their new carers did not have historical knowledge of the child i.e. past medical histories to accurately consent for immunisations or to inform healthcare professionals of the child's eligibility for vaccinations. Not having a caregiver safety net appeared to be an underlying factor in these incidents (this overarching theme of responsibility will be discussed further in the next section 3.1.2.2).

Examples

Example 29: "Baby in foster placement. Health visiting team not informed. No handover from social worker or previous health visitor. Foster carer attended clinic with baby asking for assessment of baby as not seen by health visitor since in her care. Lack of capacity in team to carry out a schedule of growing skills in a timely fashion. In this case the social worker had assumed that the foster carer would inform the HV attached to her GP that she had care of a young baby and would arranged for necessary health care e.g. routine immunisations. The social worker did not explicitly ask the foster carer to do so. She did not do so and was expecting the health visitor to contact her, when the HV did not know about the placement."

Example 30: "The nurse realised she had given a 2nd immunisation after only 18 days. Health Visitor felt bad about errors and not entering information on computer, she did follow procedure and did check the list but family had no red book for whatever reason (perhaps because they are asylum seekers)."

Example 31: "Child placed with adoptive parents. Mum advised by Social Worker to attend surgery to complete primary vaccinations. Attended surgery on with child health record, no health records available or medical records. Only two Immunisations recorded in red book, immunisation given with consent. Informed by ***** Social Services that child had already completed her primary immunisations."

3.1.2.2 Responsibility

3.1.2.2.1 Not taking responsibility/ blame

Some reports were written in a way that implied that parents, grandparents, healthcare professionals, or the children themselves were the cause of incidents. Blaming of caregivers was described in the context of them not arranging appointments in a timely manner, not holding the child appropriately during administration, forgetting to bring red books, or failing to disclose important medical information. Caregivers are typically responsible for providing school nurses with consent forms (which contain information about contraindications) via children; failing to alert school nurses to vaccine contraindications via these consent forms was described by some reports. Non-adherent children were occasionally described in a way that justified the incident. These reports reflect a culture of blame, and emphasise the importance of numerous people's roles in ensuring safe childhood vaccination and in turn the need for shared responsibility rather than blame.

Examples

Example 32: "Child given single antigen meningitis C vaccine instead of combined meningitis & Hib vaccine (Menitorix) child's records of immunisation on practice computer were confusing so nurse checked with a colleague which vaccine to give. Agreed to give menitorix but selected meningitec brand from fridge and gave this. Also grandmother who had brought child was quite anxious adding to the stress of the situation."

Example 33: "3 month old baby came for BCG immunisation - Mum assured me that BCG had not been given before, no record in child health record. Baby fulfilled criteria and BCG was given. Upon recording the immunisation on RIO it was recorded as having been given on 20th December 2011."

Example 34: "Patient is a 5 year old who has Down syndrome and is very lively. Her mother brought her to the surgery for her preschool immunisations. She did not have the red book with her, and was not appointed on the computer sheet. Due to the liveliness of the child, minimal discussion took place regarding injections and preschool booster and MMR was given. The mother was very upset when this was explained as she had not given consent for the MMR. I have informed the Dr."

Example 35: "Twins presented with their parents for immunisations, both were very distraught and it was suggested to the parents that one child had their immunisations at a time, but the children got even more distressed. The nurse went through the correct procedure and asked the relevant questions of the parents. The children were not listed on the hard copy of the schedule for pre - school boosters, but this was not uncommon, the nurse checked their medical records (the children were being very vocal and fraught at the time). The twins had received an additional immunisation."

3.1.2.2.Responsible caregivers

Several reports described caregivers advocating for their children e.g. challenging healthcare professionals decisions and actions, chasing appointments, and consequently preventing incidents from occurring, and mitigating harm to their child.

Examples

Example 36: "Mum called this office to inquire when she would receive an appointment for her baby's 3rd immunisations. She was concerned about the delay as she had been waiting since November 2009 and what consequences this would have on her child."

Example 37: "Visited patient awaiting BCG, as requested by consultant at Great Ormond Street Hospital. Parents of child have made repeated requests of a variety of professional in care of their child to find out why this has not yet been responded to."

Example 38: "Paternal Concern / Complaint raised with BCG health visitor when giving BCG at community clinic. Initially when child was born there was nobody on hospital shift to give BCG - parents were told that an appointment would be sent within a few weeks for the hospital BCG clinic. Father of child reported that his wife had phoned the hospital at least 40 - 50 times in the past 5 months trying to obtain an appointment. They did not keep a record of who they spoke to on any occasion and never heard anything back and no appointment followed. Due to travelling to India in early September they contacted their health visitor who referred to community clinic - BCG given by community service within 2 weeks of referral."

3.1.2.3 Fragmented and substandard services

3.1.2.3.1 Services at the secondary-primary care interface

Several reports described underlying system level service failures. Issues at the secondary-primary care interface were repeatedly described such as health visitors not receiving birth notifications (without these alerting them to the child's presence, vaccinations would not be organised in a timely manner). Additionally widespread communication failures between hospitals or midwives and health visitors-particularly with regards to a child's need for BCG or Hepatitis B vaccinations- underpinned several incidents.

Example

Example 39: "Audit of outcome for baby born to Hepatitis B positive mother received in child health department 6 / 7. Midwife written on bottom 'sincere apologies for not completing paperwork at delivery'. Notification received so late after delivery, very real risk that child would not have received 2nd and 3rd dose."

3.1.2.3.2 Services within primary care

Incidents involving routine childhood vaccinations were occasionally the result of miscommunication between GP surgeries/ health visitors and the child health administrative department (the public health repository of childhood vaccination records). In adolescent children poor continuity of care was described between general practice and school.

Examples

Example 40: "Mother did not receive invitation for vaccination from child health department for routine immunisations. On investigation problem with consent form being processed. Baby now missed all primary vaccinations due to this incident."

Example 41: "Telephone call from parent concerned that his child had been admitted to hospital with suspected Meningitis and this possibly could be due to not receiving an appointment for immunisation due to delay in inputting results. The results of the first vaccinations have been entered onto the child health system in order to schedule the remaining vaccinations due. Additional bank staff has been approved to deal with the backlog of inputting. Existing staff have been offered overtime hours to assist with backlog. Work priorities have been discussed with the Child Health manager to ensure that inputting of vaccinations is dealt with as an urgent priority."

3.1.2.3.3 Services on transferring between areas

Children who had moved from one locality (GP surgery) to another were also described as suffering fragmented care (records not being transferred in a timely manner, healthcare professionals in the new location not being aware of a child's presence; or difficulty registering for primary care) and vaccination-related incidents occasionally resulted from this.

Examples

Example 42: "The family informed me via an interpreter that they had moved from **** to ***** On that date I contacted the health visitor in that area and requested that she visit the family at home to complete an inward transfer visit and complete the 6 week health visitor contact. Today **** I was informed by the admin staff that the *****health visitors had returned the notes to me and they would not be visiting this child because he was not registered with their local GP. The child has therefore not received any immunisations or an 8 week development check with the GP or health visitor."

Example 43: "GP failed to advise health visiting team of transfer into area from London in ****. Not on System One, therefore 2 children not being called for routine appointments. One child missed offer of 2.5 year development review, the other not called for 12 & 13 month immunisations and 1 year development review."

Example 44: "Visited patient awaiting BCG, as requested by consultant at Great Ormond Street Hospital. Parents of child have made repeated requests of a variety of professional in care of their child to find out why this has not yet been responded to. Today I was advised by respiratory nurses that they are simply taking contact details for people requiring BCGs as there is no - one in post to provide this service. Informed previous post holder on long term sick and new recruit not yet in post."

3.1.2.3.4 Waiting lists

Substandard services were also described in the form of long and unacceptable waiting lists for newborns in need of BCG vaccine-leaving them at unnecessary risk of developing Tuberculosis.

Example

Example 45: "The baby was born to parents of mixed heritage and he will be travelling to Turkey at 4 weeks of age although eligible for BCG the baby was not referred until birth and there is a 6 week waiting list therefore he will not receive BCG before he travels."

3.1.3 Mixed methods synthesis of vaccination-related issues

This section will highlight how the themes and sub-themes identified relate to the quantitative findings, and present a visual model summarising the key weaknesses in the process of routine childhood vaccination delivery, created by combining insights from both analyses (Figure 15).

3.1.3.1 Vulnerability and associated incidents

Vulnerable children featured in a variety of incidents. The age of children made them vulnerable to certain types of incidents such as dosing errors for non-routine vaccines including: BCG, Hepatitis B, and travel vaccines. The excess doses in young children also had the potential to be larger than in older children e.g. 10 fold dosing errors were described in babies. Other incidents where the age of children made them vulnerable involved: administration of contraindicated vaccines such as flu vaccines in under 5s and HPV in pregnant teenagers; and receiving vaccines at the wrong time such those under 1 receiving the MMR - a live vaccine.

Incidents involving medically vulnerable children typically involved administration of contraindicated vaccines. However in those at risk of BCG and Hepatitis B, delays in vaccination were typical, in addition to dosing errors, administration of BCG at the wrong site or route (subcutaneously instead of intramuscularly) and administration of duplicate vaccines. These were largely because of poor coordination of care and documentation issues.

Socially vulnerable children appeared vulnerable to most vaccination-related incidents. They were vulnerable to documentation incidents (not having vaccine records or having inaccurate or out-dated records), communication failures, and poor access to primary care. Consequently they received the wrong number of doses, the wrong vaccines, and vaccines at the wrong time.

3.1.3.2 Responsibility and associated incidents

Responsible caregivers were largely described as advocating for their children when appointments had not been received i.e. vaccination was delayed, and when appointments had been sent or consent had been sought for the wrong vaccine. Not taking responsibility and blaming others was an overarching theme present in most incident types, particularly in procedural errors, communication incidents, administering the wrong vaccine, administering vaccines at the wrong time, and administering the wrong number of doses. For example, caregivers were blamed for documentation incidents i.e. 'red books' not being available, which contributed to most incident types, and not holding their child appropriately, which contributed to procedural errors. Children were also blamed for distracting health care professionals that resulted in administration of the wrong number of doses, administration of the wrong vaccine, and procedural errors.

3.1.3.3 Fragmented substandard services and associated incidents

Delayed vaccination (administration of vaccines at the wrong time) was frequently the result of: failures at the primary-secondary care interface, communication failures within primary care, failures when transferring between primary care localities, and long waiting lists. Some children received contraindicated vaccines e.g. BCG without being confirmed HIV negative as a result of failures at the primary-secondary care interface, and within primary care. Substandard services on transferring between areas typically resulted in documentation incidents, which then contributed to a variety of vaccination-related incidents.

3.1.3.4 Visual model of weakness in the vaccination process

A visual model has been created by synthesising insights from the quantitative and qualitative analysis, to display the weaknesses in the process of routine vaccination -it therefore does not apply to travel vaccines or vaccines exclusive to at-risk groups (Figure 15). Each step in the cycle represents a potential incident and example contributory factors are illustrated in yellow. At each step the child is dependent on a human or system to ensure no incidents occur. This dependency (and therefore responsibility for the child) is demonstrated at three levels: caregivers, frontline staff, and administrative system. The themes identified are over-arching and are pertinent to multiple steps in this cycle.

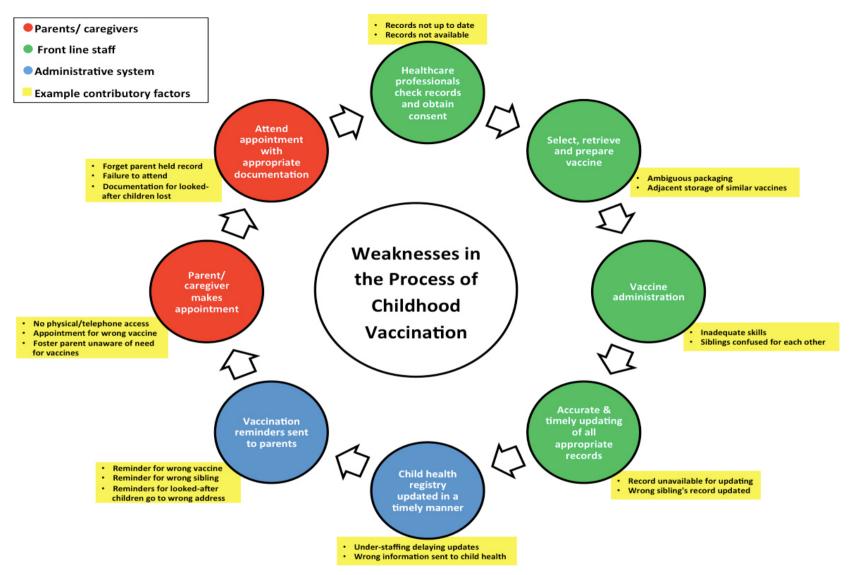


Figure 15 page 90: visual model where quantitative and qualitative insights have been combined to summarise the safety issues described by included reports

3.1.4 Summary

This analysis of vaccination-related incidents identified several priority areas requiring improvement: administration of the wrong number of doses, administration of vaccines contrary to the national vaccination schedule, and administering the wrong vaccine. To address these weaknesses, failures in documentation, and appointment management must be addressed. Factors frequently underlying these incidents were related to staff mistakes, failure to follow protocols, inadequate working conditions, and patient and caregiver factors such as behaviour, knowledge, and geography.

A key theme related to these incidents was the tendency for reporters to blame caregivers for certain failures which pre-disposed children to vaccination incidents. Also children who were medically or socially vulnerable appeared more susceptible to certain failures, which often culminated in a vaccination-related incident.

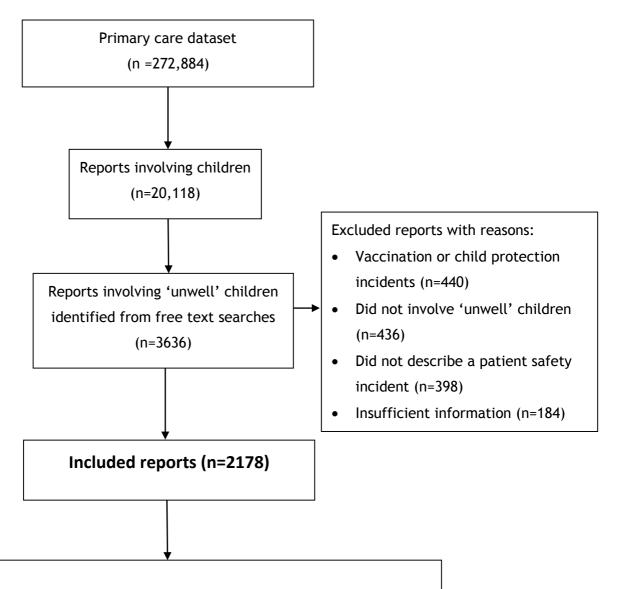
3.2 Incidents involving 'unwell' children

In this section the student will present the results of a mixed methods analysis of safety incident reports involving 'unwell' children in primary care. The student will:

- **3.2.1** provide an overview of the data included and processed, before presenting this study's key quantitative findings in more-depth
- **3.2.2** present the key themes and sub-themes identified from thematic analysis of purposively sampled reports, and describe how these themes relate to each other
- **3.2.3** explain how the qualitative insights related to the quantitative findings

3.2.1 Overview of included reports involving 'unwell' children

Of 3636 reports identified from free-text searches, 2178 were included (see Figure 16). Excluded reports: described incidents which did not involve unwell children (n=436), described incidents which only involved vaccination or child protection issues (n=440); did not describe poor care quality (n=398); and contained insufficient information (n=184). Cohen's kappa statistic for inter-rater reliability was high (k=0.72; p<0.01).



Incidents

- Total number of incidents (n=3592)
- Primary incidents (n=2191/3592)
- Contributory incidents (n=1401/3592)

Diagnoses, signs or symptoms

- Primary incidents with described diagnoses signs or symptoms (n=2032/2191)
- Number of diagnoses signs or symptoms described (n=2459)
- Number of diagnoses signs or symptoms codes (n=2582)

Contributory factors

- Primary incidents with described contributory factors (n=1219/2191)
- Number of contributory factors described (n=1785)

Outcomes

- Primary incidents with described outcomes (n=744/2191)
- Number of outcomes described (n=1004)

Figure 16 page 94: flow diagram providing an overview of how the included data were retrieved and its content

3.2.1.1 The ages of 'unwell' children involved in reported incidents

Unwell infants and pre-school children (aged less than five years old) were most frequently involved in incidents (n=1103) (see Figure 17 and Table 15). Over 25% of incidents involved unwell infants (aged <one completed year), the number of incidents reported for each subsequent age group, decreased suddenly to 124 in those aged one to two years, peaked in those aged two to three years (n=182) and decreased and plateaued between 56-106 incidents in those aged 4-18 years. Those aged 12-17 years had the highest odds of harm (Table 15).

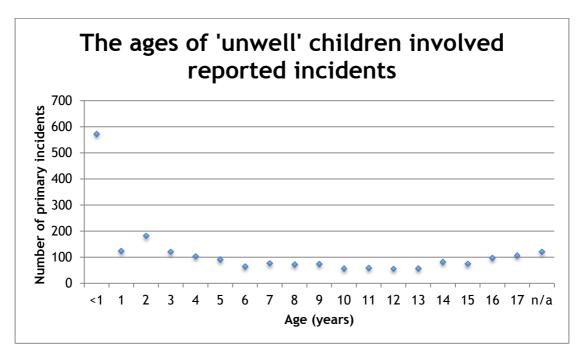


Figure 17 page 95: scatter chart illustrating the frequency and ages of 'unwell' children involved in reported incidents

Table 15 page 96: illustrates the severity of harm, the proportion of harmful incidents, and the odds of harm for each age group

Age range	Severi	ty of harm				N primary	Odds of
	No harm	Low harm	Moderate harm	Severe harm	Death	incidents (% harmful)	harm
Under 28 days	72	16	11	4	-	103 (30.1)	0.43
1 month to 1 year	360	77	43	8	3	491 (26.7)	0.36
2 to 4 years	397	85	48	8	4	542 (26.8)	0.37
5 to 11 years	378	109	48	8	3	546 (30.8)	0.44
12 to 17 years	326	100	68	13	2	509 (36)	0.56
N primary incidents	1533	387	218	41	12	2191 (30)	0.43

3.2.1.2 Temporal trends in reported incidents involving 'unwell' children

Included reports described incidents occurring between 2005-2013, the number of incidents reported increased from 49 incidents in 2005 to 438 incidents in 2011, this decreased to 299 incidents in 2013 (Figure 18).

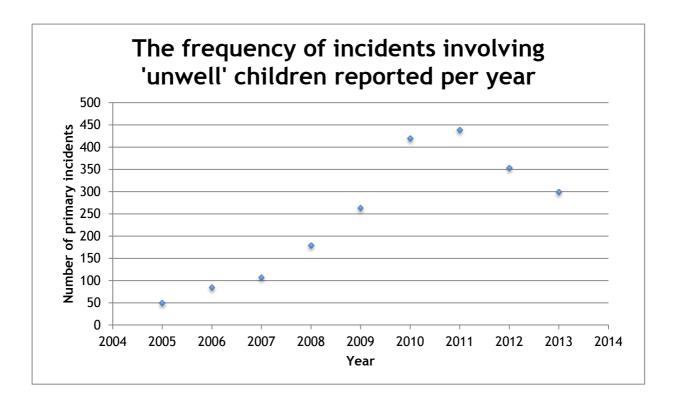


Figure 18 page 97: scatter chart illustrating the frequency of reported incidents involving 'unwell' children over time

3.2.1.3 Classification of diseases, signs or symptoms in 'unwell' children

To meet the criteria for inclusion, reports must have described an incident involving an 'unwell' child. 2032 primary incidents had associated diagnoses, signs or symptoms that enabled classification using the ICD-10, and 347 described multiple diagnoses, signs or symptoms. Of the 2032 primary incidents with associated conditions described, 1039 involved acute presentations (95 of which were in children with chronic conditions) and 699 incidents involved children with chronic conditions.

'Unwell' children experiencing safety incidents (n=1225) were frequently suffering from: respiratory disorders, non-specific signs and symptoms (such as fever), injuries, or digestive and genitourinary disorders (see Table 16). Children with digestive or genitourinary conditions, endocrine metabolic or nutrition conditions, and neurological or sensory conditions had the highest odds of harm from a reported safety incident (see Table 17).

Table 16 page 99: illustrates the conditions described in the 'unwell' children experiencing safety incidents (N.B.*some children had multiple similar conditions, signs,

Type of condition	N primary
	incidents
Respiratory system	387*
Cough, dyspnoea, tachypnoea, wheezing	127
Asthma	123
Respiratory infection	76 60
Other	69 289*
Injuries Head injuries	123
Head injuries Poisoning/ overdose accidental or of undetermined intent	42
Limb injuries	38
Burns and corrosion	28
Other	60
Non-specific signs and symptoms	281*
Fever	133
Altered consciousness, behaviour, emotions	77
Reduced food and fluid intake/ weight loss/ failure to thrive	44
Digestive and genitourinary system	268*
Disorders of oral cavity, salivary gland and jaw	74
Vomiting	69
Abdominal pain	32
Disorders of stomach, oesophagus, and duodenum	22 21
Genitourinary disorders Other	69
Skin and musculoskeletal system	245*
Rash	79
Altered skin colour	76
Other	91
Neurological and sensory system	231*
Epilepsy	126
Ear and eye disorder	61 18
Cerebral palsy and paralytic syndromes Other	34
Mental and behaviour disorders	221*
Non-specific mental health issue	65
Intentional self-harm	59
Behaviour and emotional disorders with onset in childhood and adolescence	34
Disorders of psychological development	29
Mood disorders	21
Other Infections	20 201
Non-specific infection	116
Intestinal infectious disease	49
Viral infections characterised by skin and mucous membrane lesions	12
Other	24
Endocrine, metabolic, and nutrition disorders	116*
Diabetes mellitus	72
Metabolic disorders	24
Other	21
Pregnancy, chromosomal and 'other' congenital conditions	67*
Cancer and blood	52*
Other	51
Circulatory system	50*
Total	2459

Table 17 page 100: illustrates the association between harm severity and the types of conditions present in children experiencing incidents

(N.B. * some children had multiple types of a condition; ^T some children had multiple conditions)

Type of condition	No harm	Low harm	Moderate harm	Severe harm	Death	N primary incidents (% harmful)	Odds of harm	N codes
Respiratory system	307	40*	32	7*	1	387* (20.7)	0.26	395
Non-specific signs and symptoms	216*	31*	26*	4	4*	281* (23.1)	0.30	337
Injuries	217*	36	31	4	1	289* (24.9)	0.33	291
Digestive and genitourinary system	160*	68*	28*	9	3*	268* (40.3)	0.68	287
Skin and musculoskeletal system	180*	34	24*	6	1	245* (26.5)	0.36	259
Neurological and sensory system	152*	46*	28*	4	1*	231* (34.2)	0.52	239
Mental and behaviour disorders	150*	45	19*	7	-	221* (32.1)	0.47	228
Infections	139	46	12	2	2	201 (30.8)	0.45	201
Endocrine, metabolic, and nutrition	69	23*	19	4	1	116* (40.5)	0.68	117
Pregnancy, chromosomal, other congenital	42*	15*	9*	1	-	67* (37.3)	0.60	71
Cancer and blood	38	11*	1	2	-	52* (26.9)	0.37	53
Circulatory system	34*	3	11*	2	-	50* (32)	0.47	53
Other	34	10	7	-	-	51 (33.3)	0.50	51
Number of primary incidents	1738 [™]	408 ^T	247 ^T	52 ^T	14 ^T	2459 ^T (29.3)	0.41	2582

3.2.1.4 Incident types involving 'unwell' children

The 2178 reports described 2191 primary incidents (hence the 2191 incidents referred to hereforth), and 1401 contributory incidents, therefore 3592 incidents were described in total (see Figure 16 and Appendix 3.3 for the frequencies of each combination of incidents). The most frequently described incident types involving 'unwell' children were those related to diagnosis and assessment (n=885), medication provision (n=873) adminstrative issues (n=429), communication with and about the patient (n=384), and referral for escalation of care or specialist input (n=275) (see Table 18 for the frequencies of primary incident types).

Table 18 page 102: the severity and odds of harm associated with each primary incident type

				Odds	N primary incidents		
Primary incident type I		Low harm	Moderate harm	Severe harm	Death	of harm	(% harmful)
Medication	459	143	64	6	2	0.47	674 (31.9)
Dispensing	299	69	17	1	-	0.29	386 (22.5)
Administering	75	29	18	1	-	0.64	123 (39)
Prescribing	51	12	4	1	-	0.33	68 (25)
Clinical treatment decision	26	22	14	2	2	1.54	66 (60.6)
Other	8	11	11	1	-	2.88	31 9(74.2)
Diagnosis and assessment	344	50	37	9	9	0.31	449 (23.4)
Inadequate triaging	216	13	1	-	2	0.07	232 (6.9)
Delayed assessment	65	13	9	1		0.35	88 (26.1)
Diagnosis	9	14	14	6	2	4.00	45 (80)
Insufficient assessment (non-specific)	16	5	3	-	1	0.56	25 (36)
Inadequate discharge planning	10	3	5	1	1	1.00	20 (50)
Inadequate history taking	18	1	1	-	-	0.11	20 (10)
Failure to identify high risk children	4	-	1	-	2	0.75	7 (42.9)
Inadequate examination	3	1	2			1.00	6 (50)
Other	3	-	1	1	1	1.00	6 (50)
Administrative	179	27	13	3	0	0.24	222 (19.4)
Transfer of patient information	105	16	7	-	-	0.22	128 (18)
Access to care	56	7	5	3	-	0.27	71 (21.1)
Appointment management	13	2	1	-	-	0.23	16 (18.8)
Other	5	2	-	-	-	0.40	7 (28.6)
Referral and management	135	36	32	6	1	0.56	210 (35.7)
Delayed referral	79	17	15	4	-	0.46	115 (31.3)
Failure to refer when appropriate	22	6	11	2	1	0.91	42 (47.6)
Inappropriate/ incomplete referral	24	8	5	-	-	0.54	37 (35.1)
Referral administrative issues	9	5	1	-	-	0.67	15 (40)
Failure to arrange follow up	1					0.00	1 (0)
Communication	144	20	11	2	0	0.23	177 (18.6)
Communication - patients / caregivers	127	17	10	2	-	0.23	156 (18.6)
Communication between HCPs	17	3	1	-	-	0.24	21 (19)
Treatment and procedures	53	60	26	7	-	1.75	146 (63.7)
Equipment	71	13	5	-	-	0.25	89 (20.2)
Documentation	70	2	-	-	-	0.03	72 (2.8)
Other	21	19	18	-	-	1.76	58 (63.8)
Investigations	30	11	3	-	-	0.47	44 (31.8)
Transport/ transfer of patients	27	6	9	1	-	0.59	43 (37.2)
Total	1533	387	218	41	12	0.43	2191

3.2.1.5 Factors contributing to incidents involving 'unwell' children

Of the 2191 incidents included, 1219 described at least one contributory factor and 1785 contributory factors were described in total (see Table 19 for definitions, and Appendix 3.4 for the frequency of each combination of contributory factors). Staff factors (n=722)²¹ were the most frequently described contributory factors, which included failing to follow protocols (n=356)²², mistakes (n=272)²³, and poor critical thinking (n=96). Organisational factors were also frequently described (n=463)²⁴, which included poor continuity of care (n=149)²⁵, inadequate working conditions (n=148)²⁶, and inadequate guidelines, protocols, and care plans (n=98). Other types of contributory factors described were patient (n=298)²⁷, equipment (n=78), and environmental (n=4) related. These will be discussed in further detail in relation to the incident types that they were associated with.

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²¹ In 113 cases more than one type of staff factor was described per report

²² In one report failure to follow more than 1 protocol was described

²³ In five cases more than one type of mistake was described per report

²⁴ In 84 cases more than one type of organisational factor was described per report

²⁵ In one report more than one type of continuity of care factor was described

²⁶ In 28 cases more than one type of working condition-related factor was described

²⁷ In 22 cases more than one type of patient factors was described per report

Table 19 page 104: the contributory factors described in reports

(*some reports contained descriptions of >1 type of factor)

Contributory factors - definition	N codes
Staff factors	722*
Failure to follow protocol - not adhering to organisational guidelines	356*
Mistakes - unintentional cognitive lapses	272*
Critical thinking - perception, learning, memory, concept formation, problem solving, and thin	nking 96
Knowledge - insufficient knowledge or inadequate application of knowledge	94
Other	11
Organisational factors	463*
Continuity of care - issues with the co-ordination of services	149*
Working conditions - factors relating to the work environment	148*
Inadequate protocol/ guidelines/ care plan - existing guidelines not fit for purpose	98
Education and training - insufficient education and training of staff	74
Service availability - service inaccessible to patients in a timely manner	47
Non-specific	2
Patient factors	298*
Age - child-specific factors e.g. weight-based dosing	116
Behaviour - the way in which patients or caregivers act or conduct themselves	58
Health - factors relating to the patient's physical and mental wellbeing	55
Geography - the area where patients live	38
Knowledge - insufficient knowledge or inadequate application of knowledge	30
Language - patient or caregiver unable to communicate in English	14
Looked-after- children not in the care of their parents e.g. in foster care	8
Ethnicity - the child belongs to a certain social group	1
Equipment factors - the equipment or medication is impractical, inadequate or faulty	78
Environmental factors - the physical environment is detrimental to healthcare	4
Total	1785

3.2.1.6 The severity of harm resulting from safety incidents

Of the 2191 incidents described by 2178 reports, 30% (n=658) were harmful including 12 deaths, 41 cases of severe harm, 218 cases of moderate harm, and 387 cases of low harm. Reporters only classified 19% of these incidents as harmful, 308 incidents were upgraded in terms of harm severity (see blue cells in Table 20), 25 were downgraded (see green cells in Table 20.

Table 20 page 105: the severity of harm classified by reporters and the severity described in reports using WHO definitions (*four reports describing child deaths were upgraded from moderate and severe harms)

		Reporter-allocated harm severity						
Re-coded harm	No	Low	Modera	te Seve	re Dea	th	N codes	
severity using WHO	Harm	harm	harm	harm	1			
definitions								
No harm	1521	10	1	-	1		1533	
Low harm	195	181	10	1	-		387	
Moderate harm	48	39	129	2	-		218	
Severe harm	10	2	10	19	-		41	
Death	-	-	1*	3	8		12	
N codes	1774		232 1	51	25	9	2191	

3.2.1.7 Outcomes of incidents involving 'unwell' children

Incident outcomes were described for 744 of the 2191 primary incidents; in total 1004 outcomes were described (Table 21). Patient inconvenience $(n=411)^{28}$ was the most frequently described outcome that included delays in management (n=261) and repeated visits to healthcare professionals (n=93). Clinical patient harm $(n=316)^{29}$ was also frequently described as an outcome of incidents; this included a wide range of outcomes such as harm necessitating a hospital visit (n=128), and general physical deterioration (n=101). Other types of outcomes described include psychological distress (n=59), organisational inconvenience $(n=56)^{30}$, patient injuries $(n=26)^{31}$, cardiorespiratory arrest (n=4), and death $(n=12)^{32}$

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²⁸ In 30 cases more than one type of patient inconvenience was described per report

²⁹ In 87 cases more than one type of clinical patient harm was described per report

³⁰ In one case more than one type of organisational inconvenience was described per report

³¹ In two cases more than one type of patient injury was described per report

³² In four reports death was described as a direct outcome of an incident but 12 patients who experienced incidents subsequently died

Table 21 page 107: the outcomes of incidents described in reports and their frequencies (N.B.*some reports include descriptions of multiple types of an outcome)

Outcomes	N codes
Patient inconvenience	411*
Delayed management	261
Repeated visits to healthcare providers	93
Additional treatment/ investigations	74
Unnecessary treatment	7
Other	6
Clinical patient harm	316*
Harm necessitating a hospital visit	128
General deterioration/ progression of condition	101
Discomfort/pain	27
Altered consciousness/ dizziness	25
Changes on physiological parameters	19
Nausea/ vomiting	18
Poor diabetic control	13
Seizures	12
Other	60
Psychological/ emotional distress	59
Organisational inconvenience	56*
Patient injuries	26*
Death	12
Cardio-respiratory arrest	4
Total	1004

3.2.2 Primary incident types

Almost 80% of incidents involved either medications (n=674)-, diagnosis and assessment (n=449)-, administrative(n=222)-, referral (n=210)-, or communication-related (n=177) issues. These will therefore be the focus of the exploratory quantitative analysis presented here (see Table 18 for a comprehensive breakdown of primary incident types and Table 22 for their respective contributory factors).

Table 22 page 109: the contributory factors described for each of the key primary incident types Note: *some primary incidents had multiple similar contributory factors

	Primary incident ty	pes			
Contributory factors	Medications	Diagnosis and assessment	Administrative	Referral	Communication
Staff factors	289*	205*	27*	80*	65*
Failure to follow protocol	63	140*	21	52	50
Mistakes	227*	7*	6	8	9
Critical thinking	0	63	2	21	7
Knowledge	32	23	1	15	10
Other	6	3	2	0	0
Organisational factors	148*	96*	71*	42*	27*
Continuity of care	24	25	54*	12	8
Working conditions	81*	20*	8*	9*	8
Inadequate protocols/ guidelines/ care plan	40	23	2	12	4
Education and training	17	21	3	11	6
Service availability	1	21	6	4	3
Non-specific	0	0	0	0	0
Patient factors	97*	68*	29*	28*	22*
Age	62	25	3	9	8
Behaviour	15	14	3	8	2
Health	7	15	6	4	6
Geography	5	7	14	2	2
Knowledge	15	4	1	2	4
Language	2	5	1	3	2
Looked-after	0	1	2	2	0
Ethnicity	0	0	0	1	0
Medication/ equipment factors	68	1	1	0	1
Environmental factors	0	1	0	0	0
Total number of contributory factor codes	690	423	138	177	129

3.2.2.1 Medication provision-related incidents

3.2.2.1.1 Characteristics of children experiencing medication-related incidents

The ages of children experiencing medication-related incidents crudely reflects that of the entire dataset i.e. those aged less than one year were most frequently involved in incidents, and the number of incidents per year of age plateaued between ages 2-18 years (Figure 19). Those less than 28 days old and between two to four years old had the highest odds of harm: 0.56 and 0.66 respectively; and neonates also had the highest odds of moderate harm, severe harm or death (0.27) compared to other age groups (Table 23).

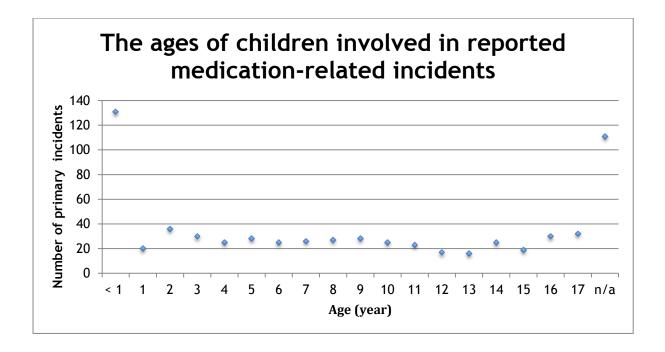


Figure 19 page 110: a scatter chart the frequency and ages of unwell children involved reported in medication-related incidents

Most (n=618) of the 674 medication-related reports described clinical conditions that necessitated treatment with medications (hence the medication-related incidents). These 618 reports described 657 diagnoses, signs or symptoms. Children with respiratory disorders (n=171) such as asthma (n=98); neurological and sensory disorders (n=113)³³ such as epilepsy (n=81); and infections necessitating antimicrobials (n=111) were most frequently involved in medication-related incidents.

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³³ In one case a child had more than one type of neurological and sensory disorder

Table 23 page 111: a cross-tabulation demonstrating the relationship between age group and harm severity

Age		S	everity of ha	rm		Proportion	Odds of	Odds of moderate harm, severe harm, or	N primary
Age	No harm	Low harm	Moderate harm	Severe harm	Death	of harm (%)	harm	death	incidents
Under 28 days	9	2	2	1	0	35.71	0.56	0.27	14
1 month to 1 year	90	20	12	2	0	27.42	0.38	0.13	124
2 to 4 years	80	36	16	1	0	39.85	0.66	0.15	133
5 to 11 years	159	49	20	1	1	30.87	0.45	0.11	230
12 to 17 years	121	36	14	1	1	30.06	0.43	0.10	173
Total	459	143	64	6	2	31.90	0.47	0.12	674

Most (n=618) of the 674 medication-related reports described clinical conditions that necessitated treatment with medications (hence the medication-related incidents). These 618 reports described 657 diagnoses, signs or symptoms. Children with respiratory disorders (n=171) such as asthma (n=98); neurological and sensory disorders (n=113)³⁴ such as epilepsy (n=81); and infections necessitating antimicrobials (n=111) were most frequently involved in medication-related incidents.

3.2.2.1.2 Primary medication-related incidents

Over 25% (n=674) of incidents were medication-related. These are broadly divided into medication dispensing- (n=386), administering- (n=123), prescribing- (n=68), and treatment decision-related incidents (n=66). (Table 18)

3.2.2.1.3 Contributory incidents

Of 674 medication-related incidents, 249 had contributory incidents and 312 contributory incidents were described in total (see Appendix 3.3 for the frequencies of each combination of incidents). The most frequent type of contributory incident was 'other medication incidents' (n=179) for example prescribing incidents often led to dispensing incidents. Other contributory incidents included communication incidents (n=39) such as giving incorrect advice to caregivers, and administrative incidents (n=30) such as inadequate transfer of patient information e.g. updated treatment plans between care settings.

3.2.2.1.4 Contributory factors

Most medication-related incidents (n=427) had contributory factors and 690 contributory factors were described in total. (Table 22 see Appendix 3.4 for the frequency of each combination of contributory factors)

3.2.2.1.5 Contributory staff factors

Staff factors $(n=289)^{35}$ were most frequently described. These included staff mistakes $(n=227)^{36}$ such as confusing similar medications (n=111), failing to follow protocols (n=63)

³⁴ In one case a child had more than one type of neurological and sensory disorder

³⁵ In 44 cases multiple staff factors contributed to medication incidents

such as double-checking procedures, and staff knowledge (n=32) for example, of certain medication contraindications particular to children.

3.2.2.1.6 Contributory organisational factors

Organisational-level contributory factors $(n=148)^{37}$ were also described by medication-related reports. These included inadequate working conditions $(n=81)^{38}$ such as inadequate staff levels and too high a workload; and inadequate guidelines protocols or care plans (40)- such as poor availability and awareness of epilepsy care plans.

3.2.2.1.7 Contributory caregiver or patient factors

Patient and caregiver level contributory factors were also frequently described (n=97)³⁹, most of these related to the patient's age (n=62) for example providing adult doses of medication, having difficulties calculating weight-based dosing, or providing medications contraindicated in certain age groups.

3.2.2.1.8 Medication or equipment-related factors

Medication and equipment-related factors were described as contributing to 68 incidents, for example storage of similar medications together resulting in the wrong medication being selected.

3.2.2.1.9 Harm and other outcomes associated with medication-related incidents

Almost a third of medication-related incidents (31.9%) were harmful, including two deaths, six cases of severe harm, 64 cases of moderate harm, and 143 cases of low harm (see Table 18). Outcomes were described for 211 incidents and 306 outcomes were described in total. These included clinical patient harm necessitating a hospital visit (n=49) (this included four children who were admitted to high dependency or intensive care); and deterioration in a child's condition (n=21). Other harmful outcomes included patient inconvenience (n=108), such as repeated visits to healthcare providers (n=52) and

³⁶ In 5 cases multiple types of mistakes contributed to medication incidents

³⁷ In 35 cases multiple organisational factors contributed to medication incidents

³⁸ In 20 cases multiple organisational factors contributed to medication incidents

³⁹ In 9 cases more than one patient/ parent factor contributed to medication incidents

delayed management of a condition (n=27); organisational inconvenience (n=19); and psychological harm (n=10).

3.2.2.1.10 Medication classes

Most medication-related incidents involved medications that target the central nervous system, the respiratory system, or infections (Table 24). Frequently reported medication types with the highest odds of harm were for: the cardiovascular system (0.75), eyes (0.63), gastrointestinal system (0.62), and infections (0.55). Medications affecting the central nervous system included anti-epileptics (n=94), anti psychotics (n=34), analgesia (n=31), and antidepressants (n=29). Respiratory medications included corticosteroids (n=89) and bronchodilators (=33). Beta lactam antibiotics (n=75) and macrolide antibiotics (n=24) were the medication types most frequently involved in infection-related medication incidents. These frequencies are likely a reflection of which medications are typically prescribed to children in primary care and their importance, and this in turn is influenced by which illnesses are frequent in childhood.

Table 24 page 115: the medications involved in medication-related incidents N.B. *some incidents involved multiple medications

Medication class	No harm	Low harm	Moderate harm	Severe harm	Death	Odds of harm	N primary incidents (% harmful)	N codes
Central nervous system	144*	38	29	3	1	0.49	215* (33)	227
Anti-epileptic	67	13	12	2	-	-	94	94
Anti-psychotic	21	4	8	1	-	-	34	34
Analgesic	21	5	5	-	-	-	31	31
Anti-depressant	19	8	2	-	-	-	29*	30
Other	15	5	2	-	1	-	23*	25
ADHD medication	12	3	-	-	-	-	15	15
Respiratory system	125*	20	12*	-	-	0.26	157* (20.4)	162
Inhaled corticosteroid	77	10	2	-	-	-	89	89
Bronchodilator	28	3	2	-	-	-	33	33
Antihistamine, immunotherapy, allergic						-		26
emergencies	14	5	7	-	-		26	
Other	10	2	2	-	-	-	14	14
Infections	97*	45*	7	1		0.55	150* (35.3)	157
Beta-lactam Beta-lactam	57	18	-	-	-	-	75*	76
Non-specific antibiotic	10	11	4	-	-	-	25*	26
Macrolide	15	7	2	-	-	-	24	24
Antiviral	11	5	-	1	-	-	17	17
Other	6	8	1	-	-	-	15	15
Endocrine system	24	6	5	-	-	0.46	35 (31.4)	36
Gastro-intestinal system	13	6	2	-	-	0.62	21 (38.1)	21
Cardiovascular system	8	2	3	1	-	0.75	14 (42.9)	14
Ear, nose, and oropharynx	9	4	-	-	-	0.44	13 (30.8)	13
Eye	8	3	2	-	-	0.63	13 (38.5)	13
Skin	12	-	1	-	-	0.08	13 (7.7)	13
Musculoskeletal and joint diseases	8	2	-	-	-	0.25	10 (20)	10
Nutrition and blood	4	4	-	-	-	1.00	8 (50)	9
Anaesthesia	1	2	4	-	-	6.00	7 (85.7)	7
Obstetrics, gynaecology and urinary-tract disorders	3	-	-	-	-	0.00	3 (0)	3
Malignant disease and immunosuppression	-	2	-	-	-	1.00	2 (50)	2
Other	1	-	-	-	1	n/a	2 (100)	2
Number of incidents	451	133	59	5	2	0.44	650 (31.1)	689

Example medication-related incidents

Example 46: "Dispensing error - prescription for Erythromycin 250mg, dispensed chlorpromazine 50mg tablets. 16-year-old patient took wrong medicine for 3 days and suffered serious side effects including catatonic seizures. Pharmacist Action - Different brand of chlorpromazine to be kept in pharmacy PCT action - contacted manufacturer to request re - assessment of packaging Similarity of packaging led to error in tablet selection."

Example 47: "GP prescribed a 5year old child chlorphenamine (antihistamine) plus 3 other items. The pharmacist dispensed chlorpromazine (anti - psychotic) instead of chlorphenamine. Mother did not recognise name so phoned pharmacy to check if it was the same. A member of staff told her that it was the same. Mother gave 8year old (sibling) 5mls of 100mg chlorpromazine. Child became extremely drowsy and was admitted to high dependency unit for observations. Child has since recovered. Pharmacy is reviewing its dispensing procedures and putting these into a written format i.e. developing standard operating procedures. Poor dispensing procedures and very limited communisation between the pharmacist and the patients."

Example 48: "Child of 8 weeks was prescribed Ranitidine 75mg / 5ml. Dose prescribed was 2.5ml twice a day. Child weighed 3.75kg. The BNF for children 2013 indicates that dose should be calculated by weight and from this it was seen that the doctor had prescribed an overdose. The dose should have been 1mg / kg three times daily. GPs checking the dose in children by weight and weighing the child accurately."

Example 49: "Chlorampenicol eye drops 0.5% were prescribed but chloramphenicol ear drops 10% were dispensed from the fridge. This occurred because the medication was dispensed in a hurry and the pharmacist did not spot the error when the second check was made. When the patient used the drops she experienced a prolonged burning sensation and was taken to the hospital when the error was recognised. The different types of chloramphenicol drops had been separated in the past and placed on different shelves due to this error occurring previously. This will now be taken further so that the ear drops are kept in enclosed containers within the fridge and clearly marked on the outside as ear drops. Similar product name. Similar package."

Example 50: "The prescription read Risperidone 1m / ml dose: 0.25mg nocte. We supplied the correct product but it was labelled 2.5ml at night. Although this is a recognised dose for a child of this age it is 10x the prescribed dose. This was a labelling error of unknown cause. The pharmacist did not pick up the labelling error. Additional care needed at time of labelling and checking, especially with children's prescriptions for unusual medications. Causes: pressure - very busy, interruptions from phone and staff."

3.2.2.2 Characteristics of children experiencing diagnosis and assessment incidents

Over half of diagnosis and assessment incidents (n=234) in 'unwell' children were in children aged less than three years (mostly in those aged under one) (see Figure 20). Children aged 12-17 had the highest odds (0.19) of more serious harm (moderate harm, severe harm or death) compared to other age groups (Table 25). Most of these 'unwell' children had: injuries (n=117)⁴⁰ such as head injuries (n=58); general signs and symptoms (n=102)⁴¹ such as fever (n=49) and altered consciousness (n=33); or skin and musculoskeletal conditions (n=87)⁴² such as rashes (n=34) and skin discolouration (n=33).

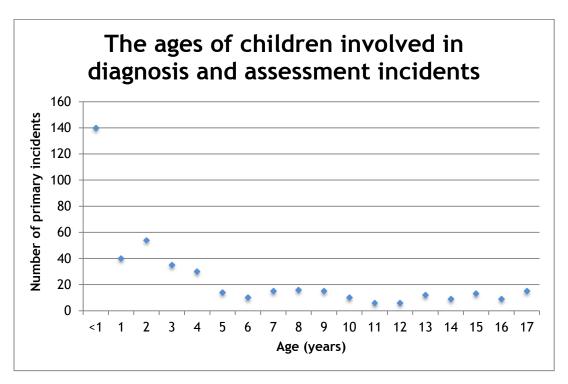


Figure 20 page 117: the frequency and ages of 'unwell' children involved in diagnosis and assessment incidents

⁴⁰ In one case multiple injuries were described

⁴¹ In 23 cases multiple non-specific signs and symptoms were described

⁴² In 12 cases multiple skin-related symptoms were described

Table 25 page 118: cross-tabulation of the relationship between age group and severity of harm

Ann			Severity of harn	Odds of	N primary			
Age	No harm	Low harm	Moderate harm	Severe harm	Death	harm	severe harm or death	incidents (% harm)
Under 28 days	6	5	1	0	0	1.00	0.09	12 (50)
1 month to 1 year	103	16	9	3	2	0.29	0.12	133 (23)
2 to 4 years	120	11	15	4	4	0.28	0.18	154 (22)
5 to 11 years	65	14	4	1	2	0.32	0.09	86 (24)
12 to 17 years	50	4	8	1	1	0.28	0.19	64 (22)
Total	344	50	37	9	9	0.31	0.14	449(23)

3.2.2.2.1 Primary diagnosis and assessment incidents

Diagnosis and assessment incidents included: insufficient assessment (n=315) (most of these (n=232) were related to inadequate patient triaging), delayed assessment (n=88), and inadequate diagnosis (n=45) i.e. delayed, missed, or wrong diagnosis.

3.2.2.2.2 Contributory incidents

Most diagnosis and assessment incidents (n=293) were preceded by other incidents: 403 contributory incidents were described in total and most were other diagnosis and assessment incidents (n=147) (see Appendix 3.3 for the frequencies of each combination of incidents). For example inadequate history taking (n=87) frequently contributed to the inadequate triaging of a child, as did failure to identify at risk and vulnerable children (n=29). Communication incidents (n=88)-particularly between healthcare professionals and caregivers (n=63)-frequently preceded incidents of inadequate diagnosis or assessment. For example, misunderstandings between triaging healthcare professionals and caregivers during telephone triaging often led to assessment issues.

3.2.2.3 Contributory factors

Most diagnosis and assessment incidents (n=302) had contributory factors and 423 were described in total (Table , see Appendix 3.4 for the frequency of each combination of contributory factors).

3.2.2.4 Contributory staff factors

Staff factors were most frequently described (n=205)⁴³. Staff failures to follow protocols were often described (n=140)⁴⁴, which included following the wrong protocol (n=96), for example assessing a child using a 'wound' protocol rather than a head injury protocol. Inadequate critical thinking (n=63) such as advising caregivers not to attend emergency care apparently in-line with protocol, despite the child being severely ill, i.e. following the protocol blindly without question, was also frequently described as contributing to these incidents.

⁴³ In 33 cases multiple staff factors contributed to diagnosis and assessment incidents

⁴⁴ In one case failure to follow more than one type of protocol contributed to a diagnosis and assessment incident

3.2.2.5 Contributory organisational factors

Organisational factors $(n=96)^{45}$ contributing to these incidents include: poor continuity of care (n=25); inadequate guidelines, protocols, or care plans (n=23); poor working conditions $(n=20)^{46}$; inadequate service provision (n=21); and insufficient education and training (n=21).

3.2.2.6 Contributory patient or caregiver factors

Patient factors $(n=68)^{47}$ contributing to these incidents included: age (n=25) such as assessing a child using an adult protocol, health (n=15) e.g. patients had pre-existing conditions that were not taken into consideration during diagnosis and assessment, and patient/ caregiver behaviour (n=14) such as non-disclosure of medical conditions.

3.2.2.2.7 Harm and other outcomes associated with diagnosis and assessment incidents

Only 23.4 % (n=105) of diagnosis and assessment incidents were harmful but they were the most harmful incident types in terms of harm severity: they resulted in 9 deaths, 9 cases of severe harm, and 37 cases of moderate harm (see Table 18). Patient inconvenience (n=104) was the most frequently described outcome type, largely because of delayed management of conditions (n=90). Other outcomes included clinical patient harm (n=54) necessitating hospital visit (n=26) and general deterioration of a child's condition (n=15).

⁴⁵ In 16 cases multiple organisational factors contributed to diagnosis and assessment incidents

⁴⁶ In two cases multiple types of working conditions contributed to diagnosis and assessment incidents

⁴⁷ In three cases multiple patient factors contributed to diagnosis and assessment incidents

Example diagnosis and assessment-reports

Example 51: "4-year-old girl was diagnosed with bilateral developmental dysplasia of the hip in a joint physiotherapy and orthopaedic clinic on. The child has been known to and seen by a number of health professionals since birth when she was diagnosed in the neonatal period as having a parietal lobe infarct with a possible risk of developing hemiplegia. There has been a significant delay in the diagnosis of this condition that may affect the outcome."

Example 52: "Patient was seen in urgent care centre with a history of passing blood in urine, treated as UTI. This was brought to my notice in the practice. I was concerned. Phoned school nurse and alerted members in practice to follow this patient. She was found to have a big tumour in her abdomen and was later diagnosed as Wilms tumour with metastasis."

Example 53: "Child with severe physical and learning disability noted to not be using her right arm in school from. She was seen in Casualty on by a Casualty Officer but not by a Paediatrician. The medical notes are not entirely legible but it appears that the examination findings showed a good range of movement in the right arm, but the impression was a pulled right elbow. Mum was told that there was a bruise on the bone and they were sent home. Child appeared to be in pain on moving the arm when visited that day. As child has complex neurodisability associated and is extremely petite on supplementary feeding but has very little subcutaneous fat my concerns were that she should have an X-ray to check for bone injury; she is also at risk of osteoporosis. I phoned the acute paediatrician; she was unaware of the child degree of disability from the casualty card but agreed for the child to attend the day unit to have a Paediatrician review and x-ray. The x-ray showed an impacted fracture at neck of humerus - child was admitted."

Example 54: "During consultation with this patient for developmental delay, I noted that patient has chest deformity. He was sent for chest X-ray. CXR report found abnormality of heart and abdominal organs- situs inversus / dextrocardia. During the second consultation I informed Dad the CXR Report and this warranted further investigations such as ultrasound of abdomen and cardiology referral for ECHO. The second line of investigation found no abnormality. Consultant radiologist advised repeat CXR. Parents were very upset on the erroneous information given to them from the CXR finding and the stress they were put through. They declined second CXR, clinic appointment and were upset with the radiographer as mother informed over the phone."

Example 55: "Patient presented to A &E with classical symptoms of new presentation of type 1 diabetes, parents concerned had presented to GP on Friday as concerned he had diabetes - GP recommended further test in 1 week later rather than immediate referral. Parents remained concerned bought blood glucose tester - sugar high. On presentation blood glucose high with 3.3 mmol / l of ketones - blood gas not acidotic. Local & national guidance of immediate referral of all suspected diabetes in children not followed."

3.2.2.3 Administrative-related incidents

3.2.2.3.1 Characteristics of children experiencing administrative-related incidents

The ages of children involved in administration-related incidents reflected that of the entire dataset, with the highest number of incidents (n=84) occurring in those aged less than 1 year (Figure 21). Most children suffering from these incidents had: general signs and symptoms (n=36)⁴⁸ such as fevers (n=13); mental and behaviour disorders (n=35)⁴⁹; respiratory conditions (n=30); and injuries (n=29).

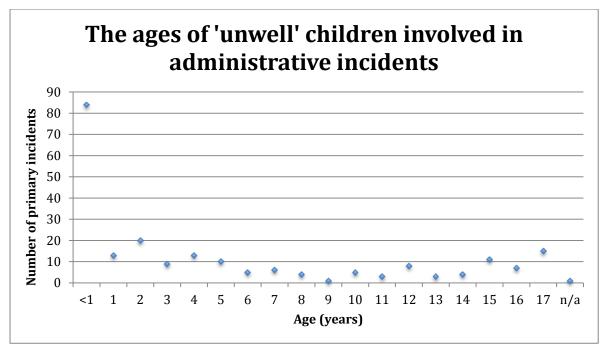


Figure 21 page 122: a scatter chart of the frequency and ages of 'unwell' children involved in administrative incidents

3.2.2.3.2 Contributory incidents

Only 100 administration-related incidents were preceded by other incidents, and these were mostly other administrative incidents (n=39) such as a patient not receiving a timely follow up phone call as a result of delays in transferring patient records; documentation failures (n=26) such as inaccurate patient records; and communication incidents (n=24) (see Appendix 3.3 for the frequencies of each combination of incidents).

⁴⁸ In seven cases multiple signs and symptoms were described

⁴⁹ In one case multiple mental and behavioural disorders were described

3.2.2.3.3 Contributory factors

Half of incidents (n=110) had contributory factors. These were mostly at the organisational level (n=71) 50 , such as poor continuity of care (n=54) 51 (see Appendix 3.4 for the frequency of each combination of contributory factors).

3.2.2.3.4 Harm and other outcomes associated with administrative incidents

Of 222 administrative incidents 19.4% were harmful which included no deaths, 3 cases of severe harm, 13 cases of moderate harm, and 27 cases of low harm. Most incidents were related to transfers of patient information between care settings (n=128) such as midwives failing to inform a health visitors about a premature baby; and access to care (n=71) such as community nurses not following up children discharged from hospital. These incidents typically resulted in inconvenience to patients/ caregivers (n=26) through delays in management (n=19) and repeated visits to healthcare professionals (n=7), and/ or clinical patient harm (n=24) for example deterioration in the child's condition (n=10).

Example administrative incidents

Example 56: "Patient with Downs syndrome was under follow up with me. Last appointment booked was for Feb 2011. It appears that this appointment was cancelled by us but no appointment was remade. Patient therefore has not had any follow up, nor screening tests recommended for children with Downs syndrome on an annual / biannual basis. This was picked up because school rang to request a report and on investigating further it appears that this child should have been under follow up. Staff have not followed procedures."

Example 57: "Deaf patient needed to be seen by an out of hours (OOH) GP. BWIC Nurse called OOH profession line several times and no reply, also called clinical shift line which was dead. Patient could not wait and was not registered with a local GP. This is a recurring theme with the new OOH GP provider, resulting in timing out of calls, inappropriate referral to A&E as unable to get through for a GP appointment, having been assessed by Nurse Practitioners."

⁵⁰ In four cases multiple organisational factors contributed to administrative incidents

⁵¹ In one case multiple continuity of care issues contributed to an administrative incident

3.2.2.4 Referral and management-related incidents

3.2.2.4.1 Characteristics of children experiencing referral-related incidents Most referral-related incidents were described in children aged less than 4 years old (Figure 22). The 'unwell' children involved in referral-related incidents mostly had non-specific signs and symptoms (n=48)⁵² including fever (n=18), altered cognition or behaviour (n=18), mental and behavioural disorders signs or symptoms (n=30) such as self-harming (n=10), and injuries (n=29), such as head injuries (n=26). Most of these children (n=165) presented acutely unwell (including 15 children with a chronic condition who presented acutely) however some referral incidents (n=19) were related to chronic condition management.

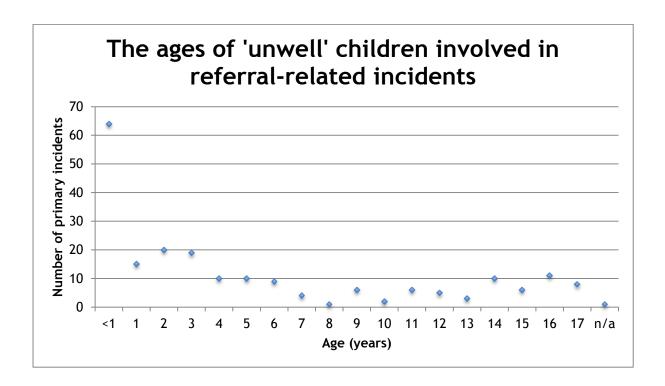


Figure 22 page 124: a scatter chart of the frequency and ages of children involved in referral-related incidents

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⁵² In 14 cases multiple non-specific signs and symptoms were described

3.2.2.4.2 Referral and management-related incident subtypes

Referral-related incidents included errors administrative referral errors such as referral paperwork getting lost or being sent to the wrong place, and errors in the referral decision-making process such as failing to refer when appropriate or making an inappropriate referral (such as referring a child with headaches to social services).

3.2.2.4.3 Contributory incidents

Most referral-related incidents (n=156) had contributory incidents and most of these were related to inadequate diagnosis and assessment (n=154) such as failing to refer a seriously unwell child as a result of not identifying the seriousness of the child's condition (see Appendix 3.3 for the frequencies of each combination of incidents). Communication-related incidents (n=30)-such as inadequate safety netting -also contributed to referral-related incidents.

3.2.2.4.4 Contributory factors

Contributory factors were described for 120 referral-related incidents (Table 22, see Appendix 3.4 for the frequency of each combination of contributory factors). Staff factors $(n=80)^{53}$ included failing to follow protocols (n=52), inadequate critical thinking (n=21), and inadequate knowledge (n=50.) Organisational level factors $(n=42)^{54}$ included inadequate protocols, guidelines or care plans (n=12), poor continuity of care (n=12), inadequate working conditions $(n=9)^{55}$, and inadequate education and training (n=11).

3.2.2.4.5 Harm and other outcomes associated with referral-related incidents

Of the 210 referral-related incidents 35.7% were harmful, which included one death, six cases of severe harm, 32 cases of moderate harm, and 36 cases of low harm (Table 18). Incident outcomes were described by half (n=105) of these reports and 125 outcomes were described. Outcomes included: patient inconvenience (n=75) such as delayed management of a condition (n=67); and clinical patient harm (n=36) such as general deterioration (n=18), harm necessitating a hospital visit (n=10), or diabetic ketoacidosis (n=4).

⁵⁴ In eight cases multiple organisational factors contributed to referral-related incidents

⁵³ In 16 cases multiple staff factors contributed to referral-related incidents

⁵⁵ In two cases multiple types of working conditions contributed to referral-related incidents

3.2.2.4.6 Example referral and management-related incidents

Example 58: "Baby admitted to A and E as SUDI aged 2 months having died at home. Baby had been seen by GP on previous evening at with temperature of 38 degrees C and possible chest infection, prescribed amoxicillin. NICE guidance for fever states that fever ≥38 in child less than 3 months is a red flag and a chi should be admitted to hospital. Preliminary results from post mortem suggesting that infection is likely cause of death.".
Example 59: "Mum attended with child alleged telephone call with health visitor at Medical Centre. Informed health visitor of non-blanching rash, glass test done and positive. Inappropriate referral to walk i centre, on arrival child referred to A&E via 999."
Example 60: "Child mother called requesting advice regarding a non-blanching rash to child upper leg also little niggly' but otherwise well, no temperature, feeding well, no sickness, bowels / bladder normal, no G in surgery therefore mother directed to walk in centre straight away for assessment. Baby should have bee referred to A&E an antibiotic injection was administered before baby was taken to A&E baby admitted for three days for observation. A viral illness was diagnosed. Urgent management, same as one above."
Example 61: "Patient reported by relative that attended centre for management of burns. Dressing applied No referral. Patient seen on 21 / 11 / 11 for a safeguarding medical, presented with full thickness burns severe enough to warrant urgent transfer and admission to a regional burns unit."

3.2.2.5 Communication incidents with or about the patient

3.2.2.5.1 Characteristics of children experiencing communication-related incidents

Communication-related incidents were most frequently reported in younger children (those aged less than three years old) (n=90) (see Figure 23). These incidents occurred in children with non-specific signs and symptoms (n=35) 56 such as fever (n=27) or failure to thrive (n=3); injuries (n=27) 57 ; gastrointestinal or genitourinary symptoms such as vomiting (n=14) and abdominal pain (n=8).

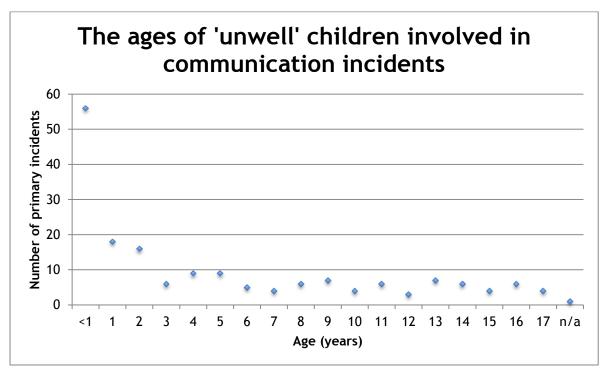


Figure 23 page 127: a scatter chart of the frequencies and ages of children involved in communication incidents

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⁵⁶ In three cases multiple non-specific signs and symptoms were described

⁵⁷ In one case multiple injuries were described

3.2.2.5.2 Contributory incidents

Most communication-related incidents (n=98, 55%) were preceded by other incidents, particularly those relating to diagnosis and assessment (n=48) e.g. giving the wrong advice to a caregiver because of a missed diagnosis, administration failures (n=11), and documentation failures (n=10) (see Appendix 3.3 for the frequencies of each combination of incidents).

3.2.2.5.3 Contributory factors

Staff factors $(n=65)^{58}$ such as failing to follow protocols (n=50) (such as safety netting protocols) were the most frequent type of contributory factor described (Table 22, see Appendix 3.4 for the frequency of each combination of contributory factors).

3.2.2.5.4 Harm and other outcomes associated with communication-related incidents

Of the 177 communicated-related incidents, 18.6% were harmful which did not include any deaths but did include 2 cases of severe harm, 11 cases of moderate harm, and 20 cases of low harm (Table 18). Most communication incidents (n=156) were between healthcare professionals and caregivers, such as inadequate safety netting, providing the wrong advice or not clearly communicating the correct advice. Most communication-related incidents (n=143, 81%) did not describe outcomes.

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⁵⁸ In 11 cases multiple staff factors contributed to communication incidents

3.2.2.5.5 Example communication incidents

Example 62: "Health Advisor sought advice on call relating to a 3 month old baby with a head injury after ending the call. Health Advisor was concerned mother would not take baby to A& E as advised. On attempting to call mother to offer nurse assessment phone was switched off. No home address details taken for caller."

Example 63: "Home visit to a child - mum very upset as she had received a letter from her child consultant detailing her most recent outpatient appointment with him. The letter stated patient condition as evolving cerebral palsy. Patient mother was extremely upset as she had never been told this as a diagnosis - it came as auite a shock to her. Patient mum has asked me to report this as an incident."

Example 64: "2nd call of the night for fever symptoms not reducing. On questioning the advice given for initial call contradicted advice to be given on return call. Relative informed that had been told to strip of clothing and tepid sponge, child was complaining of feeling cold and shivering fever remained 39.5 following medication. Concern because the same service providing contradictory advice for relief of fever symptoms. Call listened to. NA did not advise the caller to tepid sponge but did suggest that a cool flannel could be put on child's forehead but NA did advise that Ibuprofen and Paracetamol could be alternated. To be fed back to NA. Callers should not be advised to alternate Paracetamol and Ibuprofen."

Example 65: "10 year old with injury to arm, swollen and unable to move. Call was placed on queue as P3 for three hours. On attempting to call back no reply. Datix info not clear as to what the problem was with the call. HA dealt with call appropriately. The outcome was P3. The call reason was clearly documented within the call reason. The caller asked if she should take the child to A & E. The HA advised caller that she couldn't advise this but if the caller felt she wanted to then that was her choice. Caller decided to wait for a call back. Call back time was given to the caller but no worsening instructions were given. Critical thinking should have been used and clinical advice sought from the CTL on duty. HA has completed a call reflection and acknowledge she did not give worsening instructions. HA has been advised to be aware that the priority would have been higher if the this may be closer to communication category. call was triaged through limb injury."

3.2.2.6 Incidents affecting certain groups of children

The most commonly described medical conditions described in these reports will be discussed further here, including the types of incidents and harm affecting children with these conditions (see Tables 16-18).

3.2.2.6.1 Children presenting with fever

Of 133 primary incidents involving children with an unexplained fever, over 85% (n=94) were in children aged less than 5 years old. Children presenting with a fever in this dataset also had respiratory difficulty or cough (n=30), rashes (n=19), vomiting (n=19), skin discolouration (n=18), and altered consciousness or behaviour (n=14). Most incidents in children presenting with a fever involved insufficient diagnosis and assessment (n=96) such as inadequate triaging (n=50) and history taking (n=19); and communication incidents (n=51) particularly with caregivers (n=47) such as giving the wrong advice about fever management and inadequate safety netting. Two of these children died, 11 suffered moderate harm, and 6 suffered low harm.

3.2.2.6.2 Children presenting with non-specific respiratory symptoms

Of 127 primary incidents involving children with respiratory difficulty (including coughs, dyspnoea, tachypnoea, wheezing, stridor *etc.*) almost 70% (n=88) involved children aged less than 3 years. As highlighted above 30 children with respiratory difficulty had a concurrent fever, in addition 29 had skin colour changes, 14 were vomiting, 13 presented with altered consciousness, 11 had cardiac disorders or symptoms, and 8 also had a rash. In total, 248 incidents were described in these children which included: diagnosis and assessment-related incidents (n=98) such as inadequate triaging (n=45); referral-related incidents (n=33) such as failing to refer to emergency care when appropriate (n=16); and communication incidents (n=32)-particularly with caregivers (n=24). Most of these incidents did not result in harm (n=99), however 3 children suffered severe harm (which includes one child who suffered a cardio-respiratory arrest); 15 suffered moderate harm and 10 suffered low harm.

3.2.2.6.3 Children with epilepsy

The ages of the 125 children with epilepsy included in this dataset was relatively constant, ranging from <1 to <18 years old. The most frequent incident type described in these children was medication incidents (n=114), particularly dispensing (n=65) and prescribing (n=24) incidents. Diagnosis and assessment incidents (n=29) were also described in these children. Of the 125 children described in these reports, 33 suffered harm including 3 cases of severe harm (two of these children were admitted to high dependency/ intensive care, and one suffered a cardio-respiratory arrest), 15 cases of moderate harm, and 15 cases of low harm.

3.2.2.6.4 Children with asthma

Children with asthma were described by 123 reports, 102 of these related to the management of chronic asthma, 19 children presented with acute exacerbations of chronic asthma, and 2 presented with first onset/ acute asthma. Of these children 24 suffered harm: 2 severe, 5 moderate and 17 low harm. Multiple incidents (n=172) were described in these children however medication incidents were the most frequent incident type described (n=117), particularly medication dispensing errors (n=93) such as dispensing the wrong doses of corticosteroids or bronchodilators.

3.2.2.6.5 Children with head injuries

Of the 123 children with head injuries suffering patient safety incidents, 37.8% were aged less than 1 year. Most incidents related to these children involved inadequate or delayed assessment (n=122), sub-standard triaging of these children was frequently described (n=69) in addition to poor history taking (n=32). Communication incidents (n=38) were described for this group of children, such as giving poor safety netting advice to caregivers and misunderstandings between caregivers and healthcare professionals during telephone triaging. Only 12.9% of these children experienced harm, which included one death, 6 cases of moderate harm, and 8 cases of low harm. Two children with head injuries died but only one death was described as the result of healthcare failings.

3.2.3 Thematic analysis of incident reports involving 'unwell' children

Three overarching themes were identified from included reports involving 'unwell' children:

- Inadequate provision of care in the community for chronically unwell children (i.e. those requiring on-going care and/ or follow up)
- The role of caregivers in mitigating and contributing to incidents
- Weaknesses in telephone-based assessments

These will be presented, described, and supported with illustrative free-text examples; the overlap and relationship of these themes with the previously presented quantitative findings will be discussed in the mixed methods section.

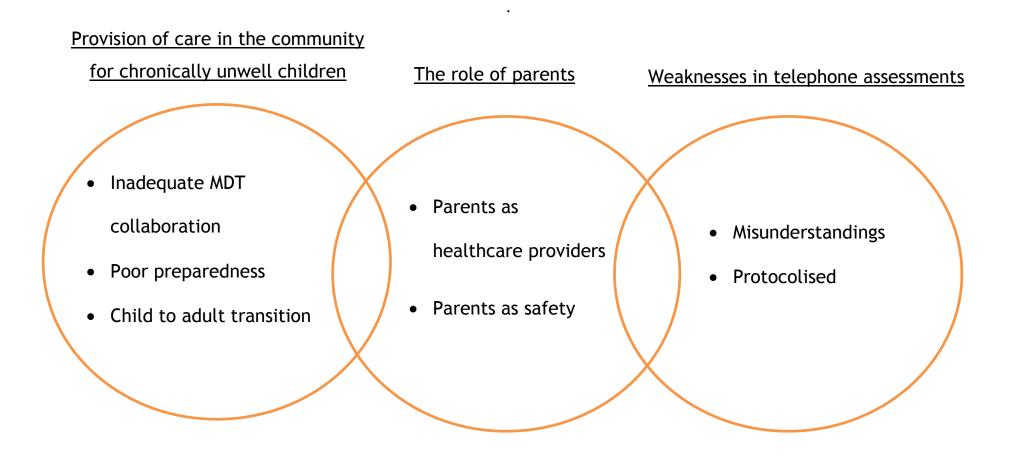


Figure 24 page 133: an illustration of the relationship between the three over-arching themes and their respective sub-themes

3.2.3.1 Provision of care in the community for complex/ chronically unwell children

3.2.3.1.1 Inadequate multidisciplinary collaboration

Poor collaboration between groups of professionals was described by reports that included health visitors, GPs, midwives, pharmacists, school nurses, social workers, and hospital clinicians. Inadequate collaboration was frequently described between healthcare professionals (hospitalists and midwives) with health visitors in the form of health visitors receiving inadequate handovers of care for neonates/ infants with jaundice, congenital disorders, birth complications, or not being updated about neonates admitted to Special Care Baby Units.

Examples

Example 66: "The mother of a baby for whom I am the Named Health Visitor, attended Baby Clinic this afternoon and informed me that her baby had been admitted to *** Hospital several weeks ago with a severe illness (treated for meningitis) and was later transferred to *** Children's Hospital where she had bowel surgery and now has an ileostomy. No liaison has been received from either hospital however the mum had been told that the Health Visitor would be informed. As a result, I have been unaware of the baby's health problems and unable to offer mum any support at a very difficult stressful time."

Example 67: "Primary visit allocated to me and I had a 'no access' visit. I left a calling card with my contact detail asking family to phone me. On return to the office, we had received a discharge letter stating baby was in SCBU. I made the decision to liaise with the staff on SCBU / midwives during the next week. Mum informed me that the baby was "starved of oxygen at birth having complex medical needs. Because of this, the family were in "conversations with children services as to whether they felt they could manage to bring the baby home". Mum was uncertain what decision she was going to make. I believe that this should have been shared with me by the hospital, as clearly, this is a complex and uncertain situation."

School nurses were also frequently not kept up-to-date about their patients' conditions, needs, and treatment plans- an issue occasionally addressed by the caregivers themselves (the only constant between the various care settings).

Example

Example 68: "Pupil had been admitted to Paediatrics following concerns about her mental health. I (school nurse) had no knowledge of this even though I have been working with the family for some time. I am concerned that both CAMHS and the children's ward failed to communicate this admission to hospital. Mum had requested support and information as to why her child had been admitted to a specialist unit. As I had very little information I felt I was unable to support Mum to the best of my capabilities."

Children with chronic conditions such as epilepsy, who should have had formal care plans known to and accessible to all relevant healthcare professionals, often either did not have these written plans in place; or they were not accessible, known of, or followed. This was particularly apparent in the pharmacy and school setting- where a unified care plan-if followed- could have prevented substandard care. In addition GPs were not always kept up-to-date by hospitalists about developments in their patients' care plan.

Example

Example 69: "8 year old child who has epilepsy seen in clinic by Paediatrician on 14 / 5. He advised changes to medication in both formulation and dose. Letter was not dictated until 28 / 6. Letter only arrived in the GP practice 16 / 7. Seen by GP on 17 / 7. In the meantime an incorrect script was issued on 5 / 7 although the parent gave the correct dose as advised by the specialist. GP phoned to mother to clarify and explain on 17 / 7 and corrected the medication."

3.2.3.1.2 Poor preparedness for care in the community

Children requiring care in the community such as those with invasive devices were often described as having issues with equipment and therapeutic adjunct provision. For example, gastrostomy feeds were often delivered in insufficient quantities, out of date, or of the wrong brand or formulation. These issues were often compounded by inadequate therapeutic adjunct provision e.g. feeding tubes. Inadequate discharge of such patients accounts for some of these inadequacies i.e. not organising community nursing, social care and adequate equipment or therapeutic adjuncts on discharge.

Example

Example 70: "Carer called to inform us that delivery from Kangaroo was incorrect. Patient hadn't received enough feed. Patient had received either too many plastics or not enough."

Failing to ensure that relevant parties in the community had adequate training to care for such children was also described i.e. discharging high-risk children without providing life-support training to caregivers; or expecting school nurses to provide care for which they have receiving no training e.g. tracheostomy care.

Examples

Example 71: "Patient is a 3 year old with cerebral palsy. She requires feeding by gastrostomy tube. She will commence school nursery on 22 / 09. It has been reported that *** is refusing to fund education / training of school staff."

Example 72: "Potential incident as training not taken place. Mum trained but CCN to visit to reinforce training and liaise with school nurse regarding: school training. 2nd home visit to reinforce training and mum updated about format school training, message left for school nurse to contact me for Epipen training. Training still hasn't taken place and care plan for school / medication still at home."

Example 73: "A child that requires oral suction was taken to school on borough transport. The escort had been trained to give oral suction. Mum told the escort that the suction was not working properly. Once on the school bus the child was sick again. Escort went to give oral suction and the pump was not working. School nurse to organise training for escort on emergency procedures. To advised if equipment faulty not to take child to school on school transport."

3.2.3.1.3 Child to adult transition

Issues with adolescents transitioning from child to adult care were evident from reports. Due to a lack of a unified definition of when an adolescent is considered an adult by the NHS, occasionally children (between 16-18 years in particular) were unable to access care as they were not classed as children by the child health service or as adults by the adult service. These issues were described for mental health services, community nursing, and the outpatient setting.

Examples

Example 74: "Patient contacted the children's community nursing Team to ask if one of the nurses would go out and give an injection that was due today. She has been discharged home with no appropriate referral and follow up care and instructions. She has also been discharged home from Adult Care services and should receive appropriate follow up with adult services as she is no longer a paediatric patient. District nurses have refused to provide any follow up care until this young lady is 18. This is the second incident that has happened to this young lady who is very sick."

Example 75: "Young adults who have been discharged from paediatric services are not being picked up by adult District Nursing Services. Young adults 16 / 17 / 18yrs are therefore at risk of having unsupported care in community unless a more formal agreement is in place to support young adults with health needs at home. No apparent service / pathway for young adults in transition phase Young adults at risk of harm if better support is not provided."

3.2.3.2 The role of caregivers

3.2.3.2.1 Caregivers as healthcare providers

In the community setting caregivers often act as healthcare providers- this was a prevalent sub-theme. Caregivers are responsible for recognising when their child is ill and deteriorating; seeking help on behalf of their child; during out-of-hours, via telephone triaging, they are responsible for assessing and triaging their child (guided by the telephone service); and they are often responsible for providing care and treatment in the community -including administering medication. Parental error- described often and permitted by the systems in place- was a key contributor to substandard care in the community. Conversely, parents were also described as preventing errors from reaching their child - which will be discussed further in the next section.

Examples

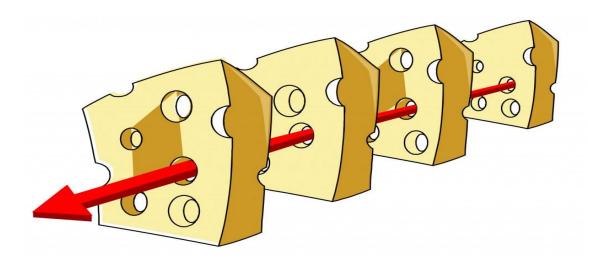
Example 76: "The parent took a copy of the prescription sheet to the GP who wrote out an FP10 prescription for the wrong dose. This was then taken to the local pharmacy and the medication was dispensed. It was the mother who noticed the error as the paediatrician had given her an instruction sheet detailing how much medication she should give."

Example 77: "Health Advisor answered' no' to a rash that looked like bleeding or bruising when the child did have a mottled purple rash making the call a P3. HA read question addressing' does she have a purple discolouration of the skin that looks like bruising or bleeding under the skin' to which the mother responded 'no'."

Example 78: "After checking the patient record I labelled the prescription. I phoned the patient mother. She had removed the labels because they weren't right. She is a nurse and understands the dosage her daughter is taking. I re-dispensed the prescription and delivered the prescription."

3.2.3.2.2 Caregivers as safety nets

Several reports described parents as 'error' safety nets e.g. using Reason's Swiss cheese model of harm, parents represented an additional layer of protection (or slice of cheese) (Reason J 1990; Walsh KE et al. 2005). Unwell children were in frequent contact with a variety of healthcare professionals responsible for different aspects of care; during and between these care encounters parents tended to be a constant presence i.e. the continuous and unifying factor (Berger Z et al. 2013). They are the expert with their own child, and engaged and informed parents were described as preventing incidents from reaching their child whereas less engaged more passive parents did not.



Examples

Example 79: "Patient had abdominal pain and vomiting, step - father went to see GP without patient for advice, GP then gave step - father prescription to collect. Prescription was for Propanolol reason not explained to step father mum read information leaflet and therefore did not give to her daughter. Felt to be an inappropriate prescription."

Example 80: "*** is prescribed 1.5mg Risperidone at 18.00hrs. The dose and time for this medication is on her medication consent form and is signed by her mother. The nursing assistant practitioner commented that she thought *** was also being given Risperidone at school at 15.00hrs. *** is only supposed to have this medication 1 x daily. A phone call was made to the nursing team who confirmed that *** was having 1.5mg Risperidone at 15.00hrs whilst she was in school. Better communication needed between school nursing teams and community nursing team. Consent forms from parents must be continuously checked with them to ensure that children are receiving the correct medicines at the correct time. It needs to be reiterated to parents that they must inform community nursing staff of any changes to their child medication."

3.2.3.3 Weaknesses in telephone-based assessments

3.2.3.3.1 Misunderstanding

Despite clear guidelines and protocols for telephone triaging of children, and robust training of healthcare professionals conducting these assessments, misunderstandings between caregivers and healthcare professionals via the telephone was a prevalent subtheme. Healthcare professionals were described as misinterpreting the information provided by callers e.g. confusing medication enquiries and symptomatic calls which required triaging for referral. Similarly callers (typically parents) were described as misinterpreting assessment questions, providing incorrect answers to assessment questions, and generally misunderstanding advice provided by healthcare professionals over the telephone.

This partly overlaps with the previous theme (the role of caregivers) - this telephone-based system is dependent upon caregivers not only assessing their child but communicating that assessment effectively. Advice about fever management appeared particularly prone to miscommunication. Caregivers were reporting that healthcare professionals had provided inappropriate advice for fever management such as tepid sponging. On review some healthcare professionals were indeed providing this inappropriate advice-but some had provided appropriate advice that was misinterpreted.

Examples

Example 81: "Call placed in Health Information Queue as a medicines call which was a potential overdose of paracetamol."

Example 82: "I have listened to the call and it was correctly assessed by the Health Advisor. The call was a 999 after nurse assessment due to the fact that the child had a non - blanching rash. However, when asked by the health advisor if the child had a rash she was advised they didn't and therefore the call was correctly prioritised as a P3."

Example 83: "4 month old baby was feverish, had one pupil larger than the other and a hard fontanelle. Call was prioritised as a P2. There was approximately a 20 minute delay before the call was then assessed by a nurse. These symptoms were all potentially very serious so I called an ambulance without any further assessment. Health Advisor used 'generally unwell' protocol, and although he asked all the questions he did use any critical thinking when the mother commented that the child was "a little bit more dazed than usual "and" drowsy not with it "and therefore entered the incorrect answer to "are they able to respond normally to you now ". HA commented that he did not know that a hard fontanelle could be dangerous."

Example 84: "During assessment of call about child with ongoing fever and diarrhoea and vomiting, mother informed me that a nurse advisor had given advice yesterday to give ibuprofen and paracetamol at 2 hourly intervals for pain relief. Call listened to. The nurse advisor gave information regarding giving ibuprofen and paracetamol, but did not say to give them together at 2 hourly intervals. Advice given by the nurse was safe."

3.2.3.3.2 Protocolised medicine

The rigid protocols for healthcare providers conducting telephone assessment were themselves described as the source of some substandard care. Numerous reports described children who had been correctly triaged using the correct protocol but in whom the outcome was not adequate i.e. the algorithm advised self-care rather than attending emergency department. To prevent re-occurrence of these incidents callers were typically advised to use 'critical thinking' and to question the protocol outcome or seek advice when an outcome seems inappropriate. This is a flaw in the system which humans are expected to detect and compensate for. These health advisors were expected to know when to deviate from protocol.

Protocols were also described as the source of error when phone calls involved more than one child, were third party i.e. a neighbour phoning on behalf of a family without a phone, where symptoms were intermittent or not accounted for by a specific protocol, and where children had multiple issues and the health advisor was unclear which issue to use as the primary issue for triaging purposes. These often resulted in children not being triaged with the most appropriate protocol or being triaged incorrectly with the correct protocol.

Example 85: "Call concerning a baby under 2 months with worsening swelling in umbilical area - baby was crying and had been unwell all day. Nurse advisor used 'other symptoms' algorithm instead of unwell baby under 3 month algorithm - she answered 2 questions and then downgraded the call from 'GP same day' to 'GP next working day'. The caller rang back a few hours later and swollen area was worsening, changing colour and baby still crying."

Example 86: "Mum reporting patient presenting with high temperature, fitting for 2 minutes and drowsiness. Patient has a history of fits. Inappropriate protocol chosen. Should have been assessed under 'fit' rather than 'fever' - as it would have covered all the correct questions and given correct end point."

3.2.4 Mixed methods synthesis of reported issues involving 'unwell children'

This section will discuss the relationship and links between the three overarching themes previously presented in section 3.2.3: provision of community care for complex patients; the role of caregivers; and weaknesses in the process of telephone assessment, and the quantitative findings reported in section 3.2.2.

3.2.4.1 Provision of care in the community and associated incidents

The sub-theme poor multidisciplinary collaboration was not associated with very harmful patient outcomes in terms of frequency or severity i.e. it was not implicated in any severe harms or deaths. Poor multidisciplinary collaboration was often associated with issues of transferring patient information between care settings (18% harmful) as well as documentation incidents e.g. medical records not being available (3% harmful). The main consequence described included fragmented knowledge about the patient between community team members. Communication incidents between healthcare professionals (19% harmful), such as inadequate handovers, were also implicated in poor multidisciplinary collaboration.

Poor preparedness for care in the community was associated with more harmful patient outcomes. This sub-theme included equipment-related incidents, which were 20% harmful though did not include any severe harms or deaths, e.g. not arranging appropriate care equipment for the home on discharge. Referral-related incidents, which were 35.7% harmful and included 7 severe harms and deaths, were associated with this sub-theme e.g. not referring to community nursing in a timely manner prior to discharge from hospital. In addition, incidents related to inadequate assessment for discharge, which were 50% harmful and included 2 severe harms and deaths, were associated with this sub-theme such as sending high-risk children home without providing caregivers with basic life support training or sending children home with an inadequate supply of medication.

This poor preparedness resulted in incidents related to treatment and procedures, which were 64% harmful and included 7 severe harms and deaths, e.g. children not receiving appropriate nursing care in the community; and medication incidents, which were 31.9% harmful including 8 severe harms and deaths, e.g. children being prescribed inadequate amounts of medications.

Issues related to transition from child to adult care was associated with several types of harmful incidents. This sub-theme was largely associated with administrative incidents such as inability to access services, which were 21% harmful. They were occasionally the result of referral incidents, which were 35.7% harmful such as adolescents being inappropriately referred to adult community nursing. However, at times they reflected a gap in services for this vulnerable age

group- this was described as resulting in treatment and procedure incidents (which were 64% harmful) e.g. not receiving necessary care.

3.2.4.2 The role of caregivers and associated incidents

This sub-theme was associated with several frequently and severely harmful incident types: diagnosis and assessment incidents (23.4% harmful which included 18 severe harms and deaths), communication with caregivers (19% harmful which included 2 severe harms), and medication incidents (31.9% harmful which included 8 severe harms and deaths).

The role of parents as healthcare providers was apparent in diagnosis and assessment incidents (particularly those related to inadequate triaging)- as they are responsible for providing accurate medical histories - and if assessment is telephone-based, parents are also responsible for assessing and examining their child.

Communication incidents between caregivers and healthcare professionals were also frequently associated with these incidents, as not only is the caregiver responsible for assessing their child, but they must also communicate that assessment, this child's history, and any concerns they may have effectively.

Parents were described as playing a prominent role in medication incidents either as partly being responsible for them, or mitigating them. In the community parents are frequently responsible for administering medications to their child- this was described as a source of error resulting in administration incidents- but it was also described as an opportunity where parents detected dispensing or prescribing errors - and therefore prevented them from reaching their child.

3.2.4.3 Weaknesses in the process of telephone-based assessment and associated incidents

Harmful incident types related to weaknesses in the process of telephone-based assessment included: diagnosis and assessment incidents which were 23.4% harmful and included 18 severe harms and deaths; and referral incidents which were 35.7% harmful and included 7 severe harms and deaths (Figure 25).

Diagnosis and assessment incidents (particularly those related to inadequate triaging and delayed assessment) were often the result of mismanaged telephone assessments. For example triaging an acutely unwell child with multiple symptoms using the wrong protocol (because the assessor is unsure which symptom to triage first e.g. fever or seizure) resulted in inadequate triaging. In addition inadequate triaging occasionally resulted in a child not being prioritised for further assessment, resulting in a delayed assessment. Referral incidents such as delayed referrals, failing to refer when appropriate and inappropriate referrals were described as the result of these assessment incidents. For example, as a result of inadequate triaging protocols, severely unwell children were referred to the out-of-hours GP service rather than the Emergency Department.

Communication incidents, particularly between callers (typically parents) and health care professionals, were prominently associated with this theme. Inadequate safety netting and failing to check callers' understanding were frequently described. Communication incidents resulted in some medication and treatment related incidents either because callers had received inappropriate treatment advice or had misunderstood appropriate advice resulting in them providing inappropriate treatment to their child-which was particularly noticeable for fever management.

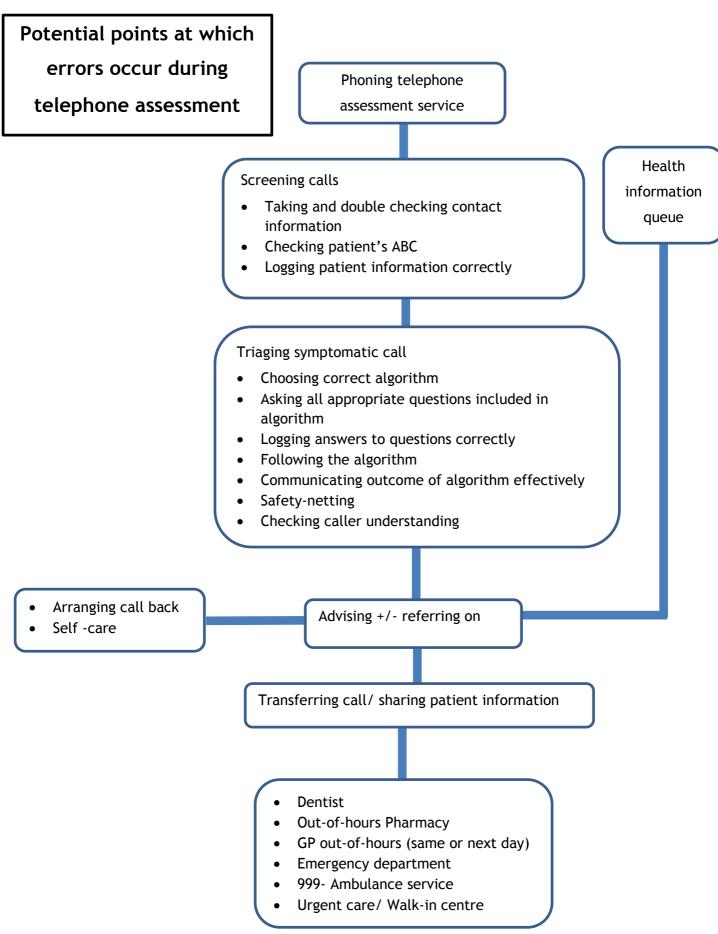


Figure 25 page 149: a visual model of the weakness in the telephone assessment system

3.2.5 Summary

Based on the burden (in terms of frequency and severity) and relative contribution of each incident type to subsequent incidents, the priority areas requiring improvement to reduce primary care-related harm to 'unwell' children include: medication-, diagnosis and assessment-, referral-, and communication-related incidents.

Children involved in medication-related incidents were largely less than one year old. These incidents were frequently related to medication dispensing, and the medications frequently involved included: anticonvulsants, bronchodilators or inhaled corticosteroids, or antimicrobials. These incidents were occasionally preceded by prescribing incidents, and were often the result of communication incidents and administrative issues. The most frequent contributory factors included staff mistakes and failure to follow protocols. These staff factors were often associated with inadequate working conditions and medication factors.

Most incidents related to diagnosis, assessment, referral, and communication were associated with telephone assessment issues, and were the result of similar contributory factors. Children involved in these incidents tended to be less than three years old, and had injuries, non-specific signs and symptoms such as fever, and skin conditions such as rashes.

A key theme, underlying many of the identified priority areas, was the contributory and sometimes protective role of parents. In the context of telephone assessments parents and caregivers were responsible for assessing the 'unwell' child and communicating their assessment effectively to the telephone advisor. Factors underlying these incidents were often related to issues with protocolised medicine including: failure to follow protocol (such as choosing the wrong assessment protocol) and failure to use critical thinking and challenge inappropriate protocol outcomes.

3.3 Literature review

The aim of this review is to identify potential and existing improvement interventions that may address the care quality issues identified in sections 3.1 and 3.2.

The research question addressed by this review is:

What improvement interventions have been proposed, tested, and/ or evaluated in primary or secondary care to address paediatric care quality issues in relation to:

- 3.3.2.2 Vaccination incidents
- 3.3.2.3 Medication incidents
- 3.3.2.4 Diagnosis and assessment incidents
- 3.3.2.5 Referral and management incidents
- 3.3.2.6 Communication incidents

3.3.1 Methods

3.3.1.1 Search strategy

A search strategy consisting of 33 keywords and Medical SubHeadings (MeSH) was designed in Medline Ovid and adapted for other databases. The search strategy was designed in three layers using terms for: the five priority areas (vaccination, mediation, diagnosis and assessment, referral, and communication) requiring improvement, child, and quality improvement (Appendix 3.3.1). The sensitivity of the search was limited to maximise specificity, for pragmatism. Searches of published literature were conducted across six databases: World of Knowledge, PsycINFO, HMIC (Health Management Information Consortium), EMBASE, Medline Ovid, and Medline in process and other non-indexed citations. Searches were limited to studies published after 2000 to ensure identified interventions were not out-dated. All references were exported to Endnote, where duplicates were removed.

3.3.1.2 Study selection

The titles and abstracts of all identified references were scanned for relevance and the full text of all those deemed potentially relevant was retrieved for further review. The student was the only reviewer involved in this process.

Studies of all designs were included: randomised controlled trials, cohort studies, cross-sectional studies, case control studies, interrupted time series, before and after studies, case series, and case reports (quality improvement reports and case studies). For inclusion: studies must have involved children aged less than 18 completed years, in primary or secondary care, and described an improvement intervention to address: vaccination-, medication-, diagnosis and assessment-, referral and management-, or communication-related incidents.

Due to the expected dearth of literature on quality improvement in primary care the student decided to include quality improvement literature from the hospital setting. Non-English studies were excluded, as well as abstracts, letters, and editorials. Studies related to increasing vaccine uptake; or to surgical, pathology or laboratory based diagnostic incidents were also excluded, as they are not relevant

to the priority areas identified as requiring improvement. Also studies from low-income countries, as defined by the World Bank in 2014, were excluded as they face different care quality challenges to the UK (World Bank 2014). Given that the purpose of this literature review was to identify improvement interventions regardless of the quality of evidence supporting them, studies were not excluded on the basis of methodological quality. However, the strength of interventions was appraised in terms of their potential to address conditions resulting in safety incidents. Strong interventions eliminate unsafe conditions by simplifying a process or removing unnecessary steps; intermediate interventions control unsafe conditions for example through developing checklists; and weak interventions accept unsafe conditions for example requiring medication double-checking.

The strength of interventions were graded using the U.S. Department of Veterans Affairs classification of strength of recommendations and these grades were corroborated by clinicians with human factors training (Morse RB and Pollack MM 2012).

3.3.1.3 Data extraction

Included studies were exported to a purpose built Microsoft Excel spread sheet, and the following variables extracted: author, year, title, study design, country, setting, priority area addressed, description of intervention(s) and the effectiveness of the intervention(s) if reported.

3.3.1.4 Data analysis

A narrative synthesis was undertaken of improvement interventions identified in relation to each of the five priority areas of interest.

3.3.2 Results

3.3.2.1 Overview of studies

Of 929 articles identified and scanned, 145 full text articles were reviewed, and 77 were included in the review (Figure 26; Appendix 3.3.3). Most were conducted in hospitals (n= 62); in the USA (n=43) or the UK (n=10); and were either interrupted time series (ITS) (n=46) or before and after studies (n=15). This review's findings are summarised by priority area in Table 26 and the characteristics of included studies are presented in Table 27.

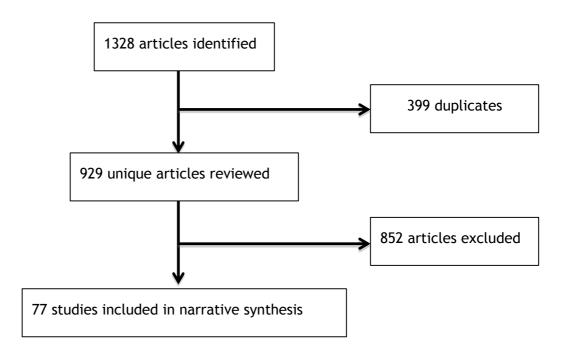


Figure 26 page 154: a flowchart demonstrating the literature review process.

Table 26 page 155: a summary of the interventions identified to address each priority area

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CU et al. 2006; Lucas AJ 2004; Mullett CJ et al. 2001; Porter SC et al. 2008; Potts AL et al. 2004; Sard BE et al. 2008; Skouroliakou M et al. 2005; Sowan AK et al. 2010; Taylor JA et al. 2008; Walsh KE et al. 2008; Warrick C et al. 2011) Bar-coding Strong (Morriss FH et al. 2009)			2009;Kazemi A et al. 2011;Kim GR et al. 2006;Kirk
2001;Porter SC et al. 2008;Potts AL et al. 2004;Sard BE et al. 2008;Skouroliakou M et al. 2005;Sowan AK et al. 2010;Taylor JA et al. 2008;Walsh KE et al. 2008;Warrick C et al. 2011) Bar-coding Strong (Morriss FH et al. 2009)			RC et al. 2005;Lehmann CU et al. 2004;Lehmann
2004; Sard BE et al. 2008; Skouroliakou M et al. 2005; Sowan AK et al. 2010; Taylor JA et al. 2008; Walsh KE et al. 2008; Warrick C et al. 2011) Bar-coding Strong (Morriss FH et al. 2009)			CU et al. 2006;Lucas AJ 2004;Mullett CJ et al.
2005;Sowan AK et al. 2010;Taylor JA et al. 2008;Walsh KE et al. 2008;Warrick C et al. 2011) Bar-coding Strong (Morriss FH et al. 2009)			2001;Porter SC et al. 2008;Potts AL et al.
2008; Walsh KE et al. 2008; Warrick C et al. 2011) Bar-coding Strong (Morriss FH et al. 2009)			2004;Sard BE et al. 2008;Skouroliakou M et al.
Bar-coding Strong (Morriss FH et al. 2009)			2005;Sowan AK et al. 2010;Taylor JA et al.
			2008; Walsh KE et al. 2008; Warrick C et al. 2011)
Pre-printed order sheets Intermediate (Alagha HZ et al. 2011: Broussard M et al.	Bar-coding	Strong	(Morriss FH et al. 2009)
/ magna / 2 of all 2011) bloadad in of all	Pre-printed order sheets	Intermediate	(Alagha HZ et al. 2011;Broussard M et al.
2009;Burmester MK et al. 2008;Cimino MA et al.			2009;Burmester MK et al. 2008;Cimino MA et al.
2004; Cunningham S et al. 2008; Kozer E et al.			2004;Cunningham S et al. 2008;Kozer E et al.
2005;Larose G et al. 2008;Robinson DL et al.			2005;Larose G et al. 2008;Robinson DL et al.
2006; Watts RG and Parsons K 2013)			2006; Watts RG and Parsons K 2013)
Environmental changes (clear Intermediate (Kaji AH et al. 2006; Sauberan JB et al.	Environmental changes (clear	Intermediate	(Kaji AH et al. 2006;Sauberan JB et al.
labelling/ storage to distinguish 2010; Sturgess E et al. 2011)	labelling/ storage to distinguish		2010;Sturgess E et al. 2011)
similar medications)	similar medications)		
Education and training Weak (Abstoss KM et al. 2011; Alagha HZ 1655al.	Education and training	Weak	(Abstoss KM et al. 2011;Alagha HZ 🖽5al.

		2011;Alemanni J et al. 2010;Bertsche T et al.
		2010;Burmester MK et al. 2008;Campino A et al.
		2009;Cimino MA et al. 2004;Davey AL et al.
		2008;Eisenhut M et al. 2011;Kozer E et al.
		2006;Leonard MS et al. 2006;Lope RJR et al.
		2009;Pallás CR et al. 2007;Robinson DL et al.
		2006;Sagy M 2009;Sullivan KM et al. 2013;Sullivan
		MM et al. 2010)
Increased pharmacist participation	Weak	(Condren M et al. 2014;Kalina M et al.
		2009;Kaushal R et al. 2008;Otero P et al.
		2008; Watts RG and Parsons K 2013)
Implementing protocols	Weak	(Sturgess E et al. 2011; Thomas C et al. 2011)
Publicising error rates	Weak	(Campino A et al. 2008;Sturgess E et al. 2011)
Verification procedures	Weak	(Watts RG and Parsons K 2013)
Diagnosis & assessment incidents		
Diagnostic decision support systems	Strong/	(Ramnarayan P et al. 2006a; Ramnarayan P et al.
	intermediate	2004;Ramnarayan P et al. 2006b)
Use clinical guidelines/ algorithms/	Intermediate	(Tabatabaei SA et al. 2012)
diagnostic checklists		
Point of care access to current	Intermediate	(Ramnarayan P et al. 2006a;Ramnarayan P et al.
evidence		2004;Ramnarayan P et al. 2006b)
Referral		
Increased patient empowerment/	Weak	(Messina FC et al. 2013)
choice		
Collaboration between referring	Weak	(Messina FC et al. 2013)
and receiving clinicians		
Communication		
Electronic handoffs linked to	Strong	(Starmer AJ et al. 2013)
medical records		
Standardised handoff templates/	Intermediate	(Kim SW et al. 2012;Sahyoun C et al. 2013;Starmer
tools/checklists		AJ et al. 2013;Weingart C et al. 2013a)
	1	
Handoff mnemonic	Intermediate	(Starmer AJ et al. 2013)
Handoff mnemonic Education and training	Intermediate Weak	(Starmer AJ et al. 2013) (Brock D et al. 2013; Hain PD et al. 2007; Starmer
		,

Table 27 page 157: a summary of the key characteristics of included studies

First author	Year	Title	Study	Hospital	Country	Priority	Interventions	Effectiveness
			type	or		area		
				primary		addressed		
				care				
Abboud	2006	Impact of	ITS	Hospital	USA	Medication	CPOE with CDS	No (no significant difference)
		workflow-						
		integrated						
		corollary orders						
		on aminoglycoside						
		monitoring in						
		children						
Abstoss	2011	Increasing	Quality	Hospital	USA	Medication	1. A poster tracking 'days since last	Yes (significant reduction in
		medication error	improve				medication error resulting in harm',	harmful medication errors (by
		reporting rates	ment				2. A continuous slideshow showing	71% p<0.01)
			report				performance metrics in the staff lounge	
							3. Multiple didactic curricula	
							4. Unit-wide e-mails summarising	
							medication errors	
							5. CPOE	
							6. Unit-based pharmacy technicians for	
							medication delivery	
							7. Patient safety report form streamlining	

Alagha	2011	Reducing	ITS	Hospital	Egypt	Medication	1.Pre-printed order sheets	Yes (significant reduction in
		prescribing errors				(prescribin	2.Education	prescribing errors p<0.001)
		in the paediatric				g)	3.Performance feedback	
		intensive care						
		unit: an						
		experience from						
		Egypt						
Alemanni	2010	An assessment of	ITS	Hospital	Canada	Medication	1.Education	n/a
		drug				(administer	2. Publicise error rates	
		administration				ing)	3.Implement protocols	
		compliance in a						
		university hospital						
		centre						

Bertsche	2010	Prospective pilot	ITS	Hospital	German	Medication	Education programme for nurses and	Yes (significant reduction in
		intervention study			у	(administer	patients	administering errors (p<0.001)
		to prevent				ing)		
		medication errors						
		in drugs						
		administered to						
		children by mouth						
		or gastric tube: a						
		programme for						
		nurses, physicians						
		and parents						
Boling	2005	Effectiveness of	ITS	Hospital	n/a	Medication	CPOE with CDS	No (non significant decrease
		computerized				(prescribin		in number of patients
		provider order				g)		requiring antidotes for
		entry with dose						prescribed opioids
		range checking on						benzodiazepines or potassium
		prescribing forms						p=0.17)

Brock	2013	Inter-professional education in team communication	Before and after study (within subject)	Hospital	USA	Communica tion	Inter-professional communication training 1.Didactic teaching session 2.Simulated session where students observe, participate, and receive feedback on communication	It changed students' attitudes towards communication skills (p<0.001), their motivation to implement them (p<0.001), and self-efficacy (p=0.005). It also changed how they perceived the utility of these skills (p<0.001).
Broussard	2009	Pre-printed order sets as a safety intervention in pediatric sedation	ITS	Hospital	USA	Medication (prescribin g)	Pre-printed order sheets	Yes (significant reduction in prescribing errors p<0.05)
Brown	2007	Error reduction when prescribing neonatal parenteral nutrition	ITS	Hospital	USA	Medication (prescribin g)	CPOE	Yes (prescription errors significantly reduced p=0.016)
Burkhart	2005	An evaluation of children's metered-dose inhaler technique for asthma medications	ITS	Ambulat ory care	n/a	Medication (administer ing)	Patient education on inhaler technique	Yes (significant reduction in administration errors p<0.001)

Burmester	2008	Interventions to	ITS	Hospital	USA	Medication	1.Pre-printed order sheets	Yes (significant reduction
		reduce				(prescribin	2.Physician education	prescribing errors (p<0.001)
		medication				g)	3. Publicising error rates	
		prescribing errors						
		in a paediatric						
		cardiac intensive						
		care unit						
Campino	2009	Educational	ITS	Hospital	Spain	Medication	1.Education	Yes (significant reduction in
		strategy to reduce				(prescribin	2.Standardised processes	prescription errors p<0.001)
		medication errors				g)	3.Updated protocols	
		in a neonatal						
		intensive care						
		unit						
Campino	2008	Medication errors	ITS	Hospital	Spain	Medication	Reviewing data and registering error rates	Yes and no (significant
		in a neonatal				(prescribin		reduction in prescribing errors
		intensive care				g and		p<0.001 and transcribing
		unit: influence of				transcribin		p=0.173)
		observation on				g)		
		the error rate						

Cimino	2004	Assessing	ITS	Hospital	USA	Medication	1.Pre-printed order sheets	Yes (significant reduction in
		medication				(prescribin	2.Real-time feedback on errors	prescribing errors p<0.001)
		prescribing errors				g)	3.Education	
		in pediatric					4.Increased pharmacist staffing	
		intensive care						
		units						
Condren	2014	Influence of a	Cross-	Outpati	USA	Medication	Pharmacist-led initiatives:	Yes (the intervention clinic
		system based	sectional	ents-		(prescribin	1. Daily prescription review	had an 11% error rate, and the
		approach to		clinic		g)	2. Provider feedback and education	comparison clinic had an error
		prescribing errors					3. EMR customisation: displayed weights in	rate of 17.4% (P<.0012))
		in a pediatric					kg only; provided recommended doses;	
		resident clinic					prepopulating instructions with dose	
							frequency and quantity; prevented selection	
							of certain formulations	
							Other changes	
							4.Education sessions on drug dosing	
							5.Provided a dosing calculator to convert	
							from mg per kg to ml (but the calculator	
							was rounding up and down to nearest	
							teaspoon)	

Cordero	2004	Impact of	ITS	Hospital	USA	Medication	CPOE with CDS	n/a (no errors in intervention
		computerized				(prescribin		group 0/117 vs. 16/136 in
		physician order				g)		comparison group)
		entry on clinical						
		practice in a						
		newborn intensive						
		care unit						
Cunningham	2008	Effect of an	RCT	Hospital	UK	Medication	1.Pre-printed order sheets	Yes (significant reduction in
		integrated care		and ED		(prescribin	2.Integrated care-pathway	prescribing errors (p=0.002)
		pathway on acute				g)		
		asthma/wheeze in						
		children attending						
		hospital: cluster						
		randomized trial						
Davey	2008	Decreasing	ITS	Hospital	UK	Medication	1.Education	No reduction in prescribing
		paediatric				(prescribin	2.Bedside prescribing guidelines	errors
		prescribing errors				g)		
		in a district						
		general hospital						

Di Pentima	2010	Impact of antimicrobial stewardship program on vancomycin use in a pediatric teaching hospital	ITS	Hospital	n/a	Medication (prescribin g)	CPOE with CDS and real-time feedback	Yes (significantly reduced patient stay p<0.05)
Dinning	2005	Chemotherapy error reduction: a multidisciplinary approach to create templated order sets	Before and after	Hospital	USA	Medication (prescribin g)	CPOE and pre-printed order sheets	Yes (prescription errors significantly reduced p<0.0001)
Eisenhut	2011	Reducing prescribing errors in paediatric patients by assessment and feedback targeted at prescribers	ITS	Hospital	UK	Medication (prescribin g)	Education	N/a (prescribing errors in intervention group 120/588 vs. comparator group 188/421)

Farrar	2003	Use of structured	ITS	n/a	n/a	Medication	CPOE with CDS	n/a (7 errors in intervention
		paediatric				(prescribin		group 7/114 vs. 46/103 in
		prescribing				g)		comparator group)
		screens to reduce						
		the risk of						
		medication errors						
		in the care of						
		children						
Fontan	2003	Medication errors	Before	Hospital	France	Medication	CPOE and unit dose drug distribution system	Yes (significant reduction in
		in hospitals:	and after			(prescribin		prescribing errors < 0.0001,
		computerized unit				g and		and administering errors
		dose drug				administeri		(p<0.001)
		dispensing system				ng)		
		versus ward stock						
		distribution						
		system						
Ginzburg	2009	Effect of a	ITS	Ambulat	USA	Medication	CPOE with CDS	Yes (significant reduction in
		weight-based		ory		(prescribin		paracetamol or ibuprofen
		prescribing				g)		prescription errors p=0.002)
		method within an						
		electronic health						
		record on						
		prescribing errors						

Gokmen-Ozel	2010	Errors in	RCT	Ambulat	UK	Medication	Use pre-measured bags of glucose	Yes (significant reduction in
		emergency feeds		ory		(dispensing		dispensing errors (p=0.03)
		in inherited)		
		metabolic						
		disorders: a						
		randomised						
		controlled trial of						
		three preparation						
		methods						
Haas-Gehres	2003	Impact of	Cross-	Primary	USA	Vaccination	Pharmacist integration in clinic	Yes significant difference in
		pharmacist	sectional	care				vaccine error rates between
		integration in a						clinics, and the intervention
		pediatric primary						clinic had significantly lower
		care clinic on						rates of missed vaccination
		vaccination errors						opportunities.(timeliness
								issues)

Hain	2007	Using risk	Cross-	Paediatr	USA	Communica	1.Crew resource management to improve	n/a
Hulli	2007	_			OJA			117 (1
		management files	sectional	ics		tion and	inter professional communication	
		to identify and		(inpatie		Medication	2.CPOE	
		address causative		nt and			3. Targeted education and feedback to	
		factors associated		outpatie			caregivers of high risk children	
		with adverse		nt)				
		events in						
		pediatrics						
Hilmas	2010	Implementation	ITS	Hospital	USA	Medication	1.CPOE with CDS	n/a (no errors in intervention
		and evaluation of				(prescribin	2.Standardised medication concentrations	group (0/200) vs. 98/200
		a comprehensive				g)	3.Education	errors in comparison group)
		system to deliver						
		pediatric						
		continuous						
		infusion						
		medications with						
		standardized						
		concentrations						

Holdsworth	2007	Impact of computerised prescriber order entry on the incidence of adverse drug	ITS	Hospital	USA	Medication	CPOE with CDS	Yes (significant reduction in total adverse drug events RR:0.64 (95%CI: 0.43-0.95))
		events in pediatric inpatients						
Hyman	2012	The use of patient pictures and verification screens to reduce CPOE errors	Before and after	Hospital and primary care	USA	Medication	Order verification screen that includes patient photograph (to reduce CPOE errors)	Yes (25% reduction in patient ID errors from n=51 in 2010 to n=37 in 2011; and a 75% reduction in the number of ordering errors)
Jani	2010	Paediatric dosing errors before and after electronic prescribing	ITS	Hospital and ambulat ory (outpati ent clinics)	n/a	Medication (prescribin g)	CPOE with CDS	Yes (significant reduction in prescribing errors p=0.001)

Kadmon	2009	Computerized order entry with limited decision support to prevent prescription errors in a PICU	ITS	Hospital	Israel	Medication (prescribin g)	CPOE with CDS	Yes (significant reduction in prescribing errors p<0.001)
Kaji	2006	Emergency medical services system changes reduce pediatric epinephrine dosing errors in the prehospital setting	ITS	Ambulat ory	USA	Medication (administer ing)	Colour coded tape for medications requiring weight-based dosing	Yes (significant reduction in administering errors 95% CI:1.4-6.6

Kalina	2009	A	ITS	Hospital	USA	Medication	MDT to care for paediatric patients	Yes (significant reduction in
		multidisciplinary		and ED		(prescribin		prescribing errors p=0.05 and
		approach to				g and		administering errors p=0.05)
		adverse drug				administeri		
		events in				ng)		
		pediatric trauma						
		patients in an						
		adult trauma						
		center						
Kaushal	2008	Unit-based	Before	Hospital	USA	Medication	increased pharmacist involvement in drug	n/a (errors in intervention
		clinical	and after				therapy	group 25/3107 vs. 45/3331 in
		pharmacists'						comparison group)
		prevention of						
		serious						
		medication errors						
		in pediatric						
		inpatients						

Kazemi	2011	The effect of	ITS	Hospital	Iran	Medication	CPOE with CDS	Yes and no (significant
		computerized				(prescribin		reduction in prescribing errors
		physician order				g and		p<0.001 but not transcribing
		entry and decision				transcribin		errors)
		support system on				g)		
		medication errors						
		in the neonatal						
		ward: experiences						
		from an Iranian						
		teaching hospital						
Kim	2012	Interdisciplinary	Evaluatio	Hospital	USA	Communica	Communication checklist and huddles	Yes- communication errors
		development and	n			tion		decreased significantly
		implementation of						
		communication						
		checklist for						
		postoperative						
		management of						
		pediatric airway						
		patients						

Kim	2006	Error reduction in	ITS	Hospital	USA	Medication	CPOE with CDS	No (163/5918 errors in
		pediatric				(prescribin		intervention group vs.
		chemotherapy:				g)		157/4978 in comparator
		computerized						group)
		order entry and						
		failure modes and						
		effects analysis						
Kirk	2005	Computer	Before	Hospital	Singapo	Medication	CPOE with CDS	Yes (significant reduction in
		calculated dose in	and after	ED and	re	(prescribin		prescribing errors (95% CI
		paediatric		ambulat		g)		0.34-0.52)
		prescribing		ory				
Kozer	2006	The effect of a	Before	ED	Canada	Medication	Education	No (no significant decrease in
		short tutorial on	and after			(prescribin		prescribing errors CI:0.66-1.7)
		the incidence of				g)		
		prescribing errors						
		in pediatric						
		emergency care						

Kozer	2005	Using a preprinted	RCT	ED	Canada	Medication	Pre-printed order sheets	Yes (significant reduction in
		order sheet to				(prescribin		prescribing errors CI:0.34-0.9)
		reduce				g)		
		prescription						
		errors in a						
		pediatric						
		emergency						
		department: a						
		randomized						
		controlled trial						
Larose	2008	Quality of orders	ITS	ED	n/a	Medication	Pre-printed order sheets	Yes (significant reduction in
		for medication in				(prescribin		prescribing errors 95% CI: 3-
		the resuscitation				g and		10% and administering errors
		room of a				administeri		95% CI: 1-6%)
		pediatric				ng)		
		emergency						
		department						
Lehmann	2004	Preventing	ITS	Hospital	USA	Medication	CPOE with CDS	Yes (significant reduction in
		provider errors:				(prescribin		prescribing errors (p<0.001)
		online total				g)		
		parenteral						
		nutrition						
		calculator						

Lehmann	2006	Decreasing errors	ITS	Hospital	USA	Medication	CPOE with CDS	n/a (prescribing errors in
		in pediatric				(prescribin		intervention group 8/142 vs.
		continuous				g)		comparator group 35/129)
		intravenous						
		infusions						
Leonard	2006	Risk reduction for	ITS	Hospital	USA	Medication	1.Education	Yes (significant reduction in
		adverse drug				(prescribin	2.Zero tolerance policy	prescribing errors p=0.001)
		events through				g)	3.Prescriber feedback	
		sequential					4. Publicise error rates	
		implementation of						
		patient safety						
		initiatives in a						
		children's hospital						
Lope	2009	A quality	ITS	Hospital	Malaysia	Medication	Education	n/a
		assurance study				(administer		
		on the				ing)		
		administration of						
		medication by						
		nurses in a						
		neonatal intensive						
		care unit						

Lucas	2004	Improving	Evaluatio	Hospital	USA	Medication	CPOE	n/a
		medication safety					- O- O-	2
		Í	"					
		in a neonatal						
		intensive care						
		unit						
MacDonald	2006	Home delivery of	Before	Ambulat	UK	Medication	Home delivery of products	Yes (significant reduction in
		dietary products	and after	ory		(dispensing		dispensing errors p<0.05)
		in inherited)		
		metabolic						
		disorders reduces						
		prescription and						
		dispensing errors						
Messina	2013	Improving	Before	Hospital	USA	Referral	New referral mechanism:	n/a (80% of those
		specialty care	and after				ED physician telephones specialist to	aged<18years kept their
		follow-up after	and areer				agree referral, accept specific treatment	appointments)
		•						арропинениз)
		and ED visit using					instructions, and agree on a referral time	
		a unique referral					frame (i.e. urgency)	
		system					2. Referral request put into electronic	
							medial record (includes time frame)	
							3. Patient chooses appointment time and	
							date (of those available in an appropriate	
							clinic within an appropriate time frame)	

Morriss	2009	Effectiveness of a	ITS	Hospital	USA	Medication	Barcode scanning	Yes (significant reduction in
		barcode						medication errors p<0.001)
		medication						
		administration						
		system in						
		reducing						
		preventable						
		adverse drug						
		events in a						
		neonatal intensive						
		care unit: a						
		prospective						
		cohort study						
Mullett	2001	Development and	ITS	Hospital	USA	Medication	CPOE with CDS	Yes (significant reduction in
		impact of a				(prescribin		prescription errors p<0.001)
		computerized				g)		
		pediatric anti-						
		infective decision						
		support program						

Neuspiel	2011	Improving reporting of outpatient pediatric medical errors	ITS	Ambulat ory care	USA	Vaccination	Eliminate distractions: 1.Use standing orders 2.Re-design room	n/a
Otero	2008	Medication errors in pediatric inpatients: prevalence and results of a prevention program	ITS	Hospital	Argenti na	Medication (prescribin g and administeri ng)	1.Increased pharmacist participation in ordering2.Education3.Reduce interruptions	Yes (significant reduction in prescription errors p<0.05 and administering errors p<0.05)
Pallas	2007	Improving the quality of medical prescriptions in neonatal units	ITS	Hospital	Spain	Medication (prescribin g)	1.Education 2.Dose calculation software	Yes (significant reduction in prescribing errors CI: 0.26-0.34)
Porter	2008	Impact of a patient-centered technology on medication errors during pediatric emergency care	Before and after	ED	USA	Medication	Parent-entered data given to provider with treatment recommendations	No (no significant reduction in medication errors p=0.42)

Potts	2004	Computerized physician order entry and medication errors in a pediatric critical care unit	ITS	Hospital	USA	Medication (prescribin g)	CPOE with CDS	Yes (significant reduction in prescription errors p<0.001)
Ramnarayan	2006	Assessment of the potential impact of a reminder system on the reduction of diagnostic errors: a quasiexperimental study	Quasi experime ntal	Hospital	UK	Diagnosis and assessment	Diagnostic reminder system- computerised decision support system (DSS)	Yes (significant decrease in diagnostic errors of omission p<0.001 (within person); and the mean diagnostic quality score increased (within person))
Ramnarayan	2004	ISABEL: a novel approach to the reduction of medical error	Descripti on	Hospital	UK	Diagnosis and assessment	Computerised decision support system (DSS)-ISABEL contains: 1.A differential diagnosis tool 2.Clinical algorithms 3.MRCPCH exam guidance 4.A section to document lessons learned from clinical errors	n/a (it was 90% accurate on evaluation)

Ramnarayan	2006	Diagnostic	Before	Hospital	UK	Diagnosis	Diagnostic reminders- computerised decision	Yes (it significantly reduced
Ť		omission errors in	and after	(paediat		and	support system (DSS)	the number of 'unsafe'
		acute paediatric	study	ric		assessment		workups p<0.001; and the
		practice: impact	(within	ambulat		assessinent		number of 'unsafe' workups
		of a reminder	subject)	ory				per case (p<0.001)
			<i>subject)</i>	•				ρει εάδε (β (0.001)
		system on		care)				
		decision-making						
Robinson	2006	Using failure	ITS	Hospital	n/a	Medication	1.Pre-printed order sheets	n/a (prescribing errors in
		mode and effects				(prescribin	2.Education	intervention group 31/221 vs.
		analysis for safe				g)	3.Policy creation	comparison group 77/331)
		administration of					4. Chemotherapy certification	
		chemotherapy to						
		hospitalized						
		children with						
		cancer						
Sagy	2009	Optimizing	ITS	Hospital	USA	Medication	Education	Yes (significant reduction in
		patient care				(prescribin		prescribing errors p<0.05)
		processes in a				g)		
		children's hospital						
		using Six Sigma						

Sahyoun	2013	Early	Evaluatio	Hospital	USA	Communica	Standard communication template	N/a- They looked at the tools
Sarryouri	2013	•		поѕрітат	USA		Standard Communication temptate	•
		identification of	n			tion		sensitivity in detecting
		children at risk						children requiring ICU not its
		for critical care						affect on errors.
Samuels	2002	Improving	Before	Primary	USA	Vaccination	1. A single person took responsibility for all	Yes- documentation errors
		accuracy in a	and after	care			vaccination-related issues	significantly decreased
		computerized					2. A clerk printed out each patients	
		immunization					computerised immunisation record before	
		registry					scheduled visits and attached it to the chart	
							(for use as a data entry form)- allowing	
							them to ID mistakes and make corrections to	
							the database	
Sard	2008	Retrospective	ITS	ED	USA	Medication	CPOE with CDS	Yes (significant reduction
		evaluation of a				(prescribin		prescribing errors p<0.001)
		computerized				g)		, , ,
		physician order				3/		
		entry adaptation						
		,						
		to prevent						
		prescribing errors						
		in a pediatric						
		emergency						
		department						

Sauberan	2010	Origins of and	Case	Hospital	USA	Medication	1.Storage solutions e.g. store paediatric and	n/a
		solutions for	studies				adult formulations separately, do not store	
		neonatal					alphabetically	
		medication-					2.Colour labelling scheme-to distinguish	
		dispensing errors					medications with similar packaging	
Skouroliakou	2005	Computer assisted	Before	Hospital	Greece	Medication	CPOE with CDS	Yes (significant reduction in
		total parenteral	and after			(prescribin		prescribing errors p<0.0001)
		nutrition for pre-				g)		
		term and sick						
		term neonates						
Sowan	2010	Computerized	Crossover	Hospital	USA	Medication	CPOE	Yes (significant reduction
		orders with						infusion order errors p<0.03)
		standardized						
		concentrations						
		decrease						
		dispensing errors						
		of continuous						
		infusion						
		medications for						
		pediatrics						

Starmer	2013	Rates of medical	Before	Hospital	USA	Communica	Handoff bundle	Yes (significantly reduced
		errors and	and after			tion	1.Communication skills training session	medical errors (p<0.001) and
		preventable					(based on TEAM STEPPS) and interactive	preventable adverse events
		adverse events					discussion session	(p=0.04)
		among					2.Mnemonic (SIGNOUT?)	
		hospitalised					3.Restructured handoffs- integrated interns'	
		children following					and senior residents' handoffs, moved to	
		implementation of					quiet and private locations, and all	
		a resident handoff					handovers were overseen by chief resident	
		bundle					or attending physician	
							4.Computerised handoff tool linked to the	
							electronic medical records	
Sullivan	2013	Personalised	Quality	Hospital	USA	Medication	Personalised performance feedback	Yes (83% reduction in
		performance	improve					pharmacist intercepted
		feedback reduces	ment					narcotic prescribing errors)
		narcotic	report					
		prescription						
		errors in a NICU						

Sullivan	2010	Impact of an	ITS	Hospital	USA	Medication	Education	Yes (significant reduction in
		interactive online				(administer		administering errors p<0.001)
		nursing				ing)		
		educational						
		module on insulin						
		errors in						
		hospitalized						
		pediatric patients						
Tabatabaei	2012	Assessment of a	Descripti	Hospital	Iran	Diagnosis	Algorithm for diagnosing and treating	n/a
		new algorithm in	on			and	children with respiratory symptoms	
		the management				assessment		
		of acute				; and		
		respiratory tract				Medication		
		infections in						
		children						

Taylor	2008	Medication	ITS	Hospital	USA	Medication	CPOE with CDS	Yes (95%CI: 0.3-0.8)
		administration				(administer		
		variances before				ing)		
		and after						
		implementation of						
		computerized						
		physician order						
		entry in a						
		neonatal intensive						
		care unit						
Thomas	2011	The impact of the	ITS	Hospital	UK	Medication	Standardising processes and updating	Yes (significant reduction in
		introduction of a				(administer	protocols	administering errors(p=0.02
		gentamicin				ing and		and monitoring errors p=0.04)
		pathway				monitoring)		
Walsh	2008	Effect of	ITS	Hospital	USA	Medication	CPOE with CDS	No (medication errors
		computer order						increased but not
		entry on						significantly)
		prevention of						
		serious						
		medication errors						
		in hospitalized						
		children						

Warrick	2011	A clinical	ITS	Hospital	UK	Medication	CPOE	Yes and no (significant
		information				(prescribin		reduction in administering
		system reduces				g and		errors (p<0.05 but no
		medication errors				administeri		prescribing errors 12/257 in
		in paediatric				ng)		intervention group vs. 14/159
		intensive care						in comparison group)
Watts	2013	Chemotherapy	Cohort	Hospital	USA	Medication	A multi-disciplinary pharmacy-associated	Yes (error rate fell by >50%)
		medication errors					error tracking system and chemotherapy	
							safety initiative	
							1.Better formatted and colour coded	
							outpatient order form	
							2.A mandatory therapy roadmap with all	
							chemotherapy orders	
							3. Double-signature on every chemotherapy	
							order	
							4. Pharmacist and nurse routinely check	
							orders prior to administration	
							5. Pharmacy standardised drug dilutions	
							6.MDT review of therapy roadmaps	
							7.CPOE (proposed)	
							8. Provide individual feedback (proposed)	

Weingart	2013	Making good	Before	Transpo	USA	Communica	Handoff bundle (education component and a	Yes (measured provider
		better	and after	rt		tion	standardised scripted handoff process)	satisfaction) p<0.05
				handoff				
				to				
				hospital				

3.3.2.2 Interventions for vaccination incidents

Childhood vaccination-related improvement was the focus of three included studies: these were all from primary care (Haas-Gehres A et al. 2014; Neuspiel DR et al. 2011; Samuels RC et al. 2002).

One article discussed improving the accuracy of vaccination-related documentation. Samuels RC et al. 2002 assessed the effectiveness of an intervention aimed at improving the accuracy of electronic vaccination records. In a before and after study they evaluated the effect of a two-pronged intervention. Firstly they allocated responsibility for all vaccination-related issues to one 'designated' person. Secondly they implemented a new process: all patients' computerised vaccination records were printed and attached to patients' charts before scheduled appointments, acting as data entry forms and allowing identification and correction of mistakes in the computerised database.

Neuspiel DR et al. 2011 propose elimination of distractions to reduce cognitive lapses i.e. mistakes which may result in vaccine administration errors. In addition Neuspiel DR et al. 2011 redesigned the vaccine room to facilitate access to the vaccination registry.

Haas-Gehres A et al. 2014 compared the effect of pharmacist integration in a paediatric primary care clinic, with a clinic that had no pharmacy service, on vaccine errors and 'missed vaccination opportunities'. Pharmacists regularly reviewed all charts and educated healthcare professionals and parents on appropriate immunisation use. Co-location of pharmacists in this manner significantly reduced the number of vaccination errors in the intervention clinic p=0.0021), and significantly reduced the number of 'missed' vaccination opportunities (p<0.001)- increasing the timeliness of vaccination.

3.3.2.3 Interventions for medication incidents

Of 64 studies focusing on reducing medication incidents, 57 were in the hospital setting.

3.3.2.3.1 Computerised physician order entry (CPOE) with or without clinical decision support (CDS)

Of the 64 studies on medication incidents, 27 described CPOE. CPOE is a term used for a computerised system used by physicians to order medications. Compared to traditional manual ordering systems, electronic systems ensure that prescriptions are complete and legible with less reliance on accurate transcription (Kaushal R et al. 2001;Rinke ML et al. 2014). CPOE systems are often combined with clinical decision support (CDS) that assesses the safety of the prescribed medications (using various algorithms) in relation to the patient's age, weight, medical history, lab results, and other prescribed medications. The CDS alerts the physician to any potential safety concerns such as interactions between medications or contraindications (Kaushal R et al. 2001).

These systems can also include forcing functions that prevent certain safety alerts from being over-ridden by physicians and ordering of certain medications e.g. preventing certain chemotherapy medications from being ordered for intrathecal administration. However CPOE systems are not error proof. Hyman D et al. 2012 tested the effect of an order verification screen-that contained patient photographs-on medication errors by aiming to improve patient identification.

Studies comparing CPOE without CDS to manual order entry reported a 44-88% reduction in prescribing error rates (Brown CL et al. 2007;Dinning C et al. 2005;Fontan JE et al. 2003;Warrick C et al. 2011) and a 21-88% reduction in administration error rates (Fontan JE et al. 2003;Warrick C et al. 2011). Whereas those evaluating the effect of CPOE and CDS on all types of medication errors in inpatients saw between a 19% increase in errors (which was not statistically significant) and a 100% reduction (Abboud PA et al. 2006;Cordero L et al. 2004;Di Pentima MC and Chan S 2010;Hilmas E et al. 2010;Jani YH et al. 2010;Kadmon G et al. 2009;Kazemi A et al. 2011;Lehmann CU et al. 2004;Lehmann CU et al. 2006;Mullett CJ et al. 2001;Potts AL et al. 2004;Rinke ML et al. 2014;Skouroliakou

M et al. 2005; Walsh KE et al. 2008). This included studies analysing all prescribing errors in inpatients, those on PICU, and those related to parenteral nutrition, anti-infective medications, and continuous infusions. Studies comparing CPOE with CDS to CPOE without CDS reported a 36-87% reduction in all prescribing errors, and specifically a 59% reduction in paracetamol and ibuprofen prescribing errors in ambulatory care (Farrar K 2003; Ginzburg R et al. 2009; Kazemi A et al. 2011; Kirk RC et al. 2005; Rinke ML et al. 2014; Sard BE et al. 2008).

3.3.2.3.2 Educational interventions

Educational interventions aimed at parents, patients, nurses, residents, and office clerks to reduce prescribing, dispensing, and administering errors were described by 17 included studies. These took the form of online tutorials, educational websites, pharmacist-led tutorials, lectures, pamphlets, practical/ simulated training sessions, and tests/ competency exams with personalised feedback on correct and incorrect answers. Alagha HZ et al. 2011 and Davey AL et al. 2008 designed point of care prescribing guidelines, i.e. dosing sheets of the most commonly used intravenous, oral and inhaled medications were available at every patient bedside.

Studies which solely evaluated the effect of educational interventions, for over three months, found an 8-87% reduction in prescribing errors, a 14-81% reduction in administering and dispensing errors and a 49-87% reduction in any type of medication error (Alemanni J et al. 2010;Bertsche T et al. 2010;Burkhart PV et al. 2005;Campino A et al. 2009;Davey AL et al. 2008;Eisenhut M et al. 2011;Kozer E et al. 2006;Leonard MS et al. 2006;Lope RJR et al. 2009;Pallás CR et al. 2007;Rinke ML et al. 2014;Sagy M 2009;Sullivan MM et al. 2010).

3.3.2.3.3 Pre-printed order sheets

Pre-printed order sheets contain formatted and pre-allocated fields that aim to decrease medication errors by increasing the completeness of hand-written medication order forms. Included studies (n=9) used pre-printed order sheets for in-patient prescribing but also specifically in relation to intensive care patients, emergency department patients, and chemotherapy prescribing. They reported a reduction in prescribing errors of between 27-82% (Alagha HZ et al. 2011; Broussard

M et al. 2009; Burmester MK et al. 2008; Cimino MA et al. 2004; Cunningham S et al. 2008; Dinning C et al. 2005; Kozer E et al. 2005; Larose G et al. 2008; Rinke ML et al. 2014; Robinson DL et al. 2006).

3.3.2.3.4 Increased pharmacist participation

Several studies (n=5) proposed increasing pharmacist participation to reduce medication errors e.g. by having ward-based pharmacists who participate in physician ward rounds and monitor drug dispensing, storage and administering. Cimino MA et al. 2004, Kaushal R et al. 2008, and Otero P et al. 2008 reported a 17-50% reduction in medication errors.

3.3.2.3.5 Medication barcoding

Barcoding of medications has been described to reduce dispensing and administering errors (Kaushal R et al. 2001). Barcoding medications enables automation of the prescription filling process and facilitates staff to rapidly match the correct medication to the correct patient during administration. For example, on scanning the medication its name, dose, route, intended patient, and staff members involved in its prescription, dispensing and administration are immediately visible (Kaushal R et al. 2001). These interventions have largely been evaluated for adult patients, however Morriss et al. 2009 reported a significant reduction (p<0.001) in all types of medication errors (ordering, transcribing, dispensing, administering, and monitoring) in the intensive care setting after implementing barcode scanning for medication administration.

3.3.2.3.6 Other interventions

Additional interventions proposed or tested by studies include: environmental changes such as changing the storage of medications or using coloured tape to distinguish similar medications; publicising error rates - to improve and maintain awareness; implementing new protocols; and verification procedures i.e. double-checking (Gokmen-Ozel H et al. 2010; Kaji AH et al. 2006; MacDonald A et al. 2006; Sturgess E et al. 2011; Thomas C et al. 2011; Watts RG and Parsons K 2013).

3.3.2.4 Interventions for diagnosis and assessment incidents

All four studies focusing on diagnosis and assessment incidents were in the hospital setting. Interventions to reduce diagnostic and assessment errors were centred on 'getting help' (Graber ML et al. 2012).

Graber et al. 2012 described 'getting help' as key to reducing cognitive burden and improving decision-making. This included having point-of-care access to current evidence-based knowledge; and using decision aids such as diagnostic checklists, diagnostic decision support systems including computerised algorithms, and clinical guidelines..

3.3.2.5 Interventions for referral incidents

One study described an intervention to improve hospital referrals (Messina FC et al. 2013). They developed a new mechanism for emergency care providers to refer on to specialists. This involved the emergency physician agreeing upon a referral plan with the specialist before discharge, noting this conversation and referral in the patient's electronic medical record, and allowing the patient to select a convenient appointment (in the agreed clinic within the agreed time frame). The aim was to improve appointment compliance, and although no figures were provided for the compliance of patients aged < 18 years before this study, post-intervention compliance was high at 80%.

3.3.2.6 Interventions for communication incidents

All seven studies proposing or evaluating interventions to improve communication focused on improving inter-professional communication (Brock D et al. 2013;Gronczewski CA 2005;Hain PD et al. 2007;Kim SW et al. 2012;Sahyoun C et al. 2013;Starmer AJ et al. 2013;Weingart C et al. 2013b). Most were hospital based (n=4) or involved the primary-hospital care interface (n=2); and most studies tested or evaluated interventions (n=6).

Educational interventions were described by Brock D et al. 2013, Starmer AJ et al. 2013 and Weingart C et al. 2013. The educational sessions implemented by Brock et al. 2013 and Starmer AJ et al. 2013 were based on the programme developed by

the Agency for Healthcare Research and Quality and the USA Department of Defence: Team Strategies, and Tools to Enhance Performance and Patient Safety (TeamSTEPPS). This approach stemmed from airlines that use crew resource management, which Hain PD et al. 2007 also implemented with an aim of reducing inter-professional communication errors (although its effect was not evaluated).

Brock et al. D 2013 evaluated the impact of a multidisciplinary educational intervention where students observed, practiced and received feedback on communication in a simulated environment, evaluating effects on student attitudes, motivation and self-efficacy towards communication skills. However Starmer AJ et al. 2013 and Weingart C et al. 2013 implemented educational interventions as part of handoff 'bundles' i.e. they tested the intervention in clinical practice rather than a simulated environment. Interventions are typically combined in a 'bundle' where it is believed that they complement each other and will be more effective together than individually.

Several studies standardised communication templates to address handoff communication errors (Kim SW et al. 2012;Sahyoun C et al. 2013;Starmer AJ et al. 2013;Weingart C et al. 2013a). Kim SW et al. 2012 implemented team 'huddles' in conjunction with a handoff checklist to ensure that patient care was handed over to the receiving team rather than from one professional to another. They saw a significant decrease in communication errors, although no specific figures were available to support this statement. Handing over to the multi-disciplinary team provided the team with an opportunity to summarise what they had understood from the handover, challenge any ambiguities, and plan for any potential patient deterioration.

Starmer AJ et al. 2013 described the most effective and comprehensive handoff bundle. It consisted of three key components: a communication training session; introduction of a handoff mnemonic (SIGNOUT?)⁵⁹ to standardise handoffs; and restructured verbal handoffs-interns and senior residents conducted handoffs together somewhere quiet, private and supervised by a chief resident or attending physician. Also, one unit linked electronic handoff documents to patient medial records. This handoff bundle significantly reduced medical errors (p<0.001) and

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⁵⁹ SIGNOUT? Sick or not for resuscitation, Identifying patient information, General hospital course, New events today, Overall health status, Upcoming possibilities, Tasks for completion, Any Questions.

preventable adverse events (p=0.04); in addition to increasing the time spent by physicians at the patient's bedside.

3.3.3 Discussion

3.3.3.1 Summary of findings

This review identified 77 studies, which described, tested, and / or evaluated a wide range of interventions to reduce: vaccination-; medication-; diagnosis and assessment-; referral-; and communication-related safety incidents.

Two studies reported a reduction in vaccination incidents by co-locating pharmacists in vaccination clinics, use of electronic vaccine records, and having a designated person responsible for vaccination issues.

Several studies reported a reduction in medication errors in response to using: CPOE and CDS; increasing pharmacist participation in medication ordering and ward rounds; pre-printed medication order sheets; and barcoding of medications to reduce medication errors.

Four studies focusing on diagnosis and assessment incidents tested or evaluated interventions such as diagnostic decision support tools. Few interventions were identified to reduce referral incidents. There was considerable support for various interventions to reduce communication errors such as: education and training, standardised handoff templates tools or checklists, and handoff mnemonics that were incorporated into a handoff bundle.

3.3.3.2 Strengths and limitations

The purpose of the literature review was to identify potential interventions that could address the priority areas requiring improvement, rather than to formally evaluate the improvement literature. Therefore, despite conducting this literature review systematically and rigorously it cannot be considered a 'systematic review'. The student was the only reviewer of identified articles and the only data extractor. In addition, the search strategy was not designed for maximal recall: its sensitivity was limited to maximise precision.

Evaluations of improvement interventions are often not to the standard of traditional effectiveness studies. In addition, as a result of broad inclusion criteria, included studies were heterogeneous in terms of their design, population, and aims.

The implications of this review's findings in relation to creating recommendations for practice and further research will be discussed in Chapter 5.

Chapter 4: Discussion of methods

4.1 Summary

Broadly this study aimed to address two questions:

- What paediatric safety incidents in primary care, involving vaccination, and the management of 'unwell' children, are occurring in practice and are reported to the NRLS?
- How can safety problems involving paediatric vaccination and the management of 'unwell' children be addressed?

The student sought to address these questions by analysing incident report data. This involved designing a search strategy for maximum recall of relevant reports; systematically capturing the content described within the free-text elements of reports; capturing the nuanced and context-specific issues present but not explicitly described within reports; and subsequently analysing this large 'processed' data set to identify learning.

4.2 Searching the NRLS data

The search strategy was dependent on reports specifying a patient age of <18 years and containing at least one key word permutation. The list of key words was developed to be as sensitive as possible but within practical limits for numbers identified. Searches were conducted in Microsoft Excel, however the specificity of searches could have been enhanced if software permitting the use of search strings containing Boolean operators had been available. Unfortunately, a notable number of reports (n=139,847; 51.2%) some of which likely involved children, did not specify patient age, and therefore would not have been retrieved for free-text searching. In hindsight the number of paediatric-related reports identified for free-text searching could have been increased with free-text searches to identify child-specific characteristics, and through using manual filters to identify additional reports in child-specific locations such as schools.

4.3 Data processing

During the student's previous pilot work with the NRLS dataset it became apparent that existing taxonomies used for classifying patient safety incidents were inadequate (Rees P et al. 2015). The level of detail of other taxonomies was insufficient and they were not grounded in UK primary care data, which limited their utility. The student therefore developed three taxonomies to describe incident types, contributory factors, and incident outcomes.

These taxonomies were developed iteratively: they were regularly amended and updated as more reports were read and different learning emerged, and are therefore grounded in UK paediatric-related primary care data. However, existing taxonomies, such as Learning from International Networks about Errors and Understanding Safety in Primary Care (LINNEAUS Euro-PC) and the WHO International Classification for Patient Safety, did at a high level, inform the taxonomies used in this study (World Health Organisation 2009). For example, the student used the "five rights" of medication administration: right medication, right patient, right dose, right route, and right time, but supplemented it with more detailed codes such as right formulation and right number of doses (Kron T 1962). This similarity will facilitate comparisons between previous and future studies in this area.

To structure the application of codes from the taxonomies, the student used the Recursive Model of Incident Analysis developed by the Australian Patient Safety Foundation (Hibbert PD et al. 2007). This allows the incident, the factors leading to the incident (i.e. other incidents and contributory factors), and the outcomes of the incident to be captured systematically, by coding these factors in chronological order. Following training, this model ensures reproducibility between coders and allows the rich detailed descriptions to be coded, rather than reduced and summarised. This model permits detailed data processing, unlike other models of incident analysis that traditionally find one 'best fit' category in which to place the incident, even if multiple incidents are described. Also, other models tend not to clearly distinguish between contributory factors and incidents (Hibbert PD et al. 2007).

During data processing the student noted all medications involved in incidents and where applicable the signs, symptoms or diagnoses present in children involved in

safety incidents. Capturing this supplemental information provided additional structured variable data, for a more in-depth analysis of the data than previous studies of this nature: as the pre-allocated medication fields are often incomplete, and signs, symptoms, and diagnoses, are not routinely captured by incident reporting systems. Using ICD-10 to classify the 'types of illness' present in children involved in incidents will also permit corroboration of this study's findings with future primary care case note reviews (the funding for which has recently been secured by the Division of Population Medicine, Cardiff University, in collaboration with Nottingham University et al.).

The NRLS does not collect information on patient disease; this information is therefore only present if reporters include it in the free-text descriptions of incidents. Therefore, the reports included in this study may not be a true reflection of all reported incidents involving 'unwell' children in primary care. Additionally this likely compromised the comprehensiveness and accuracy at which illnesses described by included reports could be classified using ICD-10. This may also be responsible for the notably high number of included incidents involving acutely unwell children. The student hypothesises that incidents involving acutely unwell children may be more likely to contain descriptions of 'illness', permitting detection during free-text searches and classification with ICD-10, than incidents involving children with chronic conditions.

4.4 Data analysis

Using the Recursive Model of Incident Analysis resulted in a complex and in-depth characterisation of reports. This allowed modelling of the complex sequence of events leading to and resulting from an incident. In turn this allowed me to examine the sequences of 'codes' and the relationships between variables using frequency distributions and cross-tabulations. To the student's knowledge few studies have analysed the events described within incident report data in this level of detail. This exploratory approach to quantitative analysis is arguably a more flexible and comprehensive approach to analysing incident report data, allowing hypotheses to be generated, rather than rigidly testing pre-formed hypotheses as done in other studies (Alexander DC et al. 2009;Bundy DG et al. 2008;Bundy DG et al. 2009;Rinke ML et al. 2010).

4.5 Thematic analysis

Supplementing quantitative analysis with thematic analysis of a purposive sample of reports provided greater insight than data processing could have alone. It enabled examination of contextual insights that would otherwise have gone unnoted. However, in retrospect it was possible to capture most of the information contained within reports using the structured coding frameworks.

In previous work with NRLS data the student had not systematically captured variables such as 'medications involved in incidents' and 'diagnoses'. Thematic analysis (aimed at examining factors not detected by the frameworks) was a useful method of exploring and not overlooking these important factors. However during this study, the scope of thematic analysis to identify 'new' contextual insights was limited as the quantitative analysis was very detailed.

4.6 Reflection on alternative approaches and methods

If the student were to do the study again, she would have done it in three parts: a case note review, incident report data analysis, and supplementary root cause analysis.

Analysis of incident report data provided in-depth learning about the nature of unsafe care in practice sufficient to address the research question. However, this study's findings are unlikely to be representative of what occurs in practice, hence the need for a case note review.

To conduct a case note review, the student would first (through consensus methods) have piloted a UK-specific paediatric trigger tool for primary care. A case note review would allow me to gain a handle on the true priority areas in primary care requiring improvement, as the findings would be representative of what occurs in practice and would also therefore be generalisable. More in-depth information could be retrieved using this method, permitting more reliable, comprehensive, and accurate classification of patient outcomes including harm severity, and patient characteristics including social status, presentation, and pre-existing diagnoses. This approach would permit identification of the most harmful incidents in primary care (in terms of frequency and severity). Also, this approach

would enable risk modelling, whereby the association between patient characteristics and safety incidents or healthcare harm could be analysed. Case note reviews have well-publicised attributes - they provide generalisable and comprehensive information about 'what' occurs in practice. However, they are unable to provide in-depth information on contributory factors, 'why' safety incidents are occurring in practice, hence the need for incident report analysis. Analysis of incident report data, as done in this study, would be necessary, to address the priority areas identified through case note review and to design interventions to mitigate particularly harmful safety incidents.

A key and unique attribute of incident report data is its ability to provide vast quantities of comprehensive and relatively detailed information on the contributory factors pre-disposing to safety incidents, as perceived by reporters. Analysing incident report data allows identification of patterns of incidents and contributory factors underlying safety incidents, which can be targeted by improvement interventions.

Unfortunately reporters do not routinely receive human factors or root cause analysis training. Therefore they are unlikely to comprehensively detect and report the system factors and root causes of incidents, they are more likely to report human-factors that are more readily apparent to them. To overcome this selective deposit bias and to supplement the information gained from incident report data, the student would conduct a root cause analysis of a sample of the most complex and harmful safety incidents. This would ensure that system issues underlying safety incidents are not overlooked and are adequately targeted by interventions.

Chapter 5: Discussion of findings

This chapter will summarise and discuss key findings related to: vaccination failures presented in section 3.1, and diagnosis, assessment, referral, and communication failures involving 'unwell' children presented in section 3.2, in the context of current literature. Recommendations for improvement in these areas, informed by the interventions identified in section 3.3, will be presented. This chapter will conclude with recommendations for future research to improve understanding of the problems identified by this study, and to further inform changes to policy and practice to improve primary care for children.

5.1 Discussion of vaccination-related findings

5.1.1 Summary

Most reported vaccination-related incidents were in younger children especially those aged less than one year old. The most frequently reported vaccination incidents were related to administration, including the wrong number of doses, wrong timing, and wrong vaccine. Other frequently reported incidents involved adverse reactions to vaccines and communication failures with parents and caregivers. The reported reasons for these incidents included documentation failures, appointment management problems, staff mistakes such as confusing similar vaccines, staff failing to follow protocols such as preparing more than one vaccine concurrently, and patient/ caregiver factors such as inappropriate behaviour and ill-health. Parents and caregivers played an important role in contributing to and preventing vaccination-related incidents, and vulnerable children such those in care (without parents to advocate for them) appeared disproportionately vulnerable to vaccination-related incidents.

5.1.2 Context of current literature

The frequency of reports describing administration of the wrong number of doses is unsurprising, because they are errors of commission i.e. as a result of actions taken, and are therefore typically apparent to the healthcare professionals involved, and therefore likely to be reported (World Health Organisation 2009). This finding also mirrors the literature, which highlights that receiving additional vaccines is a prevalent issue and often the result of inadequate documentation (Bundy DG et al. 2009;Feikema SM et al. 2000;Weltermann BM et al. 2014). A study conducted by Feikema SM et al. 2000, highlights that 20% of children in the USA receive unnecessary duplicate vaccinations.

The study contained numerous reports of delayed vaccination or receipt of vaccines out-of-sync with the national vaccination schedule. The consequences of delayed vaccination are well acknowledged in the literature, a Confidential Enquiry of child deaths related to primary care identified vaccination timeliness as a priority area requiring improvement (Derrough TF and Kitchin NR 2002; Harnden A et al. 2009; Vivier PM et al. 1999). Errors of omission, as result of actions not taken, are more difficult to detect and less likely to be reported, therefore the three child deaths detected in the study may represent a much larger body of children

who have not received vaccinations and thus vulnerable to life-threatening diseases (World Health Organisation 2009). The potentially harmful consequences of deviating from the national vaccination schedule, which is specifically developed by experts to afford children maximum protection and to minimise the risk of vaccine interactions, are unclear (Bundy DG et al. 2009; Derrough TF and Kitchin NR 2002; Vivier PM et al. 1999).

Administration of the wrong vaccine is a widespread and well-recognised problem (Bundy DG et al. 2009;National Patient Safety Agency 2004;National Patient Safety Agency 2008;Weltermann BM et al. 2014). Other studies in this area report staff confusing vaccines with similar names or packaging, as an important cause of these errors, which mirrors this study's findings (Bundy DG et al. 2009;Makeham MAB et al. 2004;National Patient Safety Agency 2004). Fortunately in the student's study no cases of severe harm or death were described as a result of administration of the wrong vaccine. However these data potentially reflect the tip of the iceberg, and of note, WHO guidelines have highlighted several case reports of severe harm or death in children who received the wrong vaccination (World Health Organisation 2014). Of concern, are children who are under-protected and whose inadequate immunity is also unrecognised, for example they received the wrong vaccine but the error went undetected.

Other studies highlight the potentially severe consequences of vaccination incidents involving live vaccines and medically vulnerable children, such as those present in the student's study. For example, a 2007 case report describes an immunocompromised child who develops severe varicella after receiving a live varicella vaccine (which was contraindicated) (Jean-Philippe P et al. 2007).

Socially vulnerable children appeared disproportionately prone to reported vaccination-related incidents, implying that the inverse care law, where those most in need of high quality care are the least likely to receive it, may be an existing problem in this context (Hart JT 1971). Vaccination uptake in socially vulnerable children is well described as sub-optimal in the literature (Barnes P et al. 2005; Webb E et al. 2001; Williams J et al. 2001). However the vulnerability of these children to vaccination-related incidents, such as receiving the wrong vaccine, is not well-publicised, this may partly be due to the difficult nature of conducting research in marginalised populations (Webb E 2004).

5.1.3 Recommendations for improvement

Recommendations to improve vaccination in children are centred on four underlying and contributory weaknesses: reducing staff mistakes; minimising documentation and appointment failures; improving caregiver knowledge; and improving staff knowledge.

Manufacturing-targeted recommendations include encouraging manufacturers to create vaccines with different packaging and names, or at least to continue using tall man lettering with the aim of reducing staff mistakes (Filik R et al. 2006;Levine SR et al. 2001).

At a policy level, electronic red books that are accessible to caregivers and staff could reduce documentation discrepancies by eliminating the current system, where three types of vaccination records exist, and staff would not be dependent on caregivers to bring red books to appointments. This could prove particularly beneficial to looked-after children who were frequently without red books in this study. CDS software could be incorporated into these electronic books, whereby warnings about vaccines contraindicated for each child appear on the screen acting as a reminder for healthcare professionals and caregivers.

Education-level recommendations to reduce incidents in medically vulnerable children include providing staff with feedback on the learning generated from vaccination-related reports raised locally, and educating caregivers and staff about vaccine contraindications and the potentially fatal consequences of receiving contraindicated vaccines. Additionally, encouraging caregiver involvement, and creating a culture where caregivers feel comfortable challenging healthcare professionals, could prevent safety incidents. At a practice level, to reduce staff mistakes, staff must adhere to verification and standardised preparation procedures.

5.2 Discussion of medication-related findings

5.2.1 Summary

Children aged less than one year typically experienced medication incidents. These children were being treated for epilepsy, asthma, and various infections. Most incidents were related to dispensing or administering of medications, and prescribing incidents often preceded them. Reasons for these incidents included communication failures between staff or with patients or caregivers, and administrative (paper-work) issues. Underlying staff factors included failure to follow protocols, mistakes, and inadequate knowledge. Patient and caregiver factors included age-specific factors (such as weight-based dosing), poor knowledge, and inappropriate behaviour. These were often compounded by medication factors such as adjacent storage of similar medications, and organisational factors such as busy working conditions or inadequate care plans for treatment of chronic conditions.

5.2.2 Context of current literature

Medication errors are believed to be the most common type of medical error in children and adults (Department of Health 2000; Department of Health 2001; Kohn LT et al. 1999; Wong IC et al. 2009). Avery AJ et al. 2013 report that children under 15 years old have an 87% excess risk of a prescribing error in UK general practice, compared to those aged 15-64, and others report that medication errors are three times more prevalent in children than adults (Kaushal R et al. 2001; Wong IC et al. 2009). Therefore this study's finding, that medication incidents are the most frequent type of reported safety incident involving 'unwell' children, is in keeping with the literature.

The ages of children involved in reported medication incidents was as expected, as similar trends are reported in the literature where younger children are more prone to medication errors (Koren G et al. 1986;Koren G and Haslam RH 1994;Smith MD et al. 2014;Wong IC et al. 2009). A national study of medication errors in young children in the USA reports that children aged <1 year are responsible for 25.2% of reported incidents in those aged less than six years (Smith MD et al. 2014). In addition, Smith MD et al. 2014 report that these younger children are also more likely to suffer severe harm from medication incidents, which is reflective of this study's findings. They hypothesise that younger children's limited ability to communicate may account for the inverse association between error rate and age.

This study's findings partially reflect those reported by Smith MD et al. 2014 where reported out-of-hospital medication errors in children aged < 6 years in the USA most frequently involved analgesics, cough and cold medications, antihistamines, and antimicrobials. However, anti-epileptic and asthma medications were the most frequent types of medications involved in medication errors in this study. Differences could be the result of international variation, differences in study populations, or unsystematic reporting bias present in both studies. Antimicrobials are one the most frequently reported class of medications responsible for adverse reactions resulting in hospital admission, and are reported as the class of medications most frequently involved in medication errors (Ghaleb MA et al. 2006;Smyth RM et al. 2012). This may be partly due to how frequently they are prescribed in practice (Ghaleb MA et al. 2006).

A recent study of medication prescribing errors in UK general practice report considerably different findings in terms of the risk of error associated with medication class (Avery AJ et al. 2013). Avery et al. 2013 report that patients (children and adults) were most at risk of error if they were prescribed musculoskeletal, malignancy, immunology/ vaccines, or skin medications. Medications were classified using the British National Formulary (as done for the student's study) however their population included adults and children, which, in conjunction with reporting bias, may explain the differences in the medication types involved in incidents (Avery AJ et al. 2013). In addition Avery AJ et al. 2013 had a narrower focus, prescribing errors in general practice, compared to this study, which focused on all medication errors reported in primary care.

The high frequency of asthma-related medication incidents was anticipated since asthma is the most common chronic condition of childhood in the UK, and a recent report highlights severe failures in the management of childhood asthma in the community (Asthma UK 2015). Asthma UK estimate that approximately 12,000 children with asthma experienced a prescribing incident between 2010-2013, and that approximately 2000 children have been prescribed long acting beta agonists alone (without inhaled steroids) putting them at risk of a severe asthma attack. Whilst the student's study's findings reflect some of these widely reported issues, it also highlights the importance of dispensing errors, which are less widely reported in this group of children.

Several studies highlight the importance of clear communication with caregivers and patients to prevent medication errors (Stebbing C et al. 2007; Wong IC et al. 2006; Wong IC et al. 2009). Walsh KE et al. 2013 highlight the contribution of communication failures with parents to home-based medication errors. In the community setting, where parents are responsible for understanding their child's treatment plan and administering the correct medication, at the correct dose, via the correct route, and at the correct time, suboptimal communication about this process increases vulnerability to medication errors (Walsh KE et al. 2011; Walsh KE et al. 2013).

A UK epilepsy review found that poor communication with parents and caregivers prompted numerous medication administration errors (Royal College of Paediatrics and Child Health 2013). Typical examples include dosing errors as a result of: confusing mg and ml and confusing different formulations of buccal midazolam (Royal College of Paediatrics and Child Health 2013). Other examples in the literature of communication failures resulting in medication errors include: failing to explain treatment plan changes and to provide written management plans for caregivers, resulting in them continuing to administer medications according to an old treatment regime. In addition, failure to communicate the importance of regularly checking expiry dates of infrequently used *pro re nata* (PRN) medications, is described as resulting in administration of expired medications (Walsh KE et al. 2011).

The national epilepsy review found that only 77% of children with epilepsy had an emergency plan for seizure management, and only 26% had a school health plan (Royal College of Paediatrics and Child Health 2013). Failing to provide a written management plan also predisposes to communication errors between caregivers, which have been reported as resulting in medication administration errors (Walsh KE et al. 2011).

The UK epilepsy review also highlights a lack of evidence of collaboration with young patients in current practice (Royal College of Paediatrics and Child Health 2013). This study highlights the importance of involving caregivers and patients in discussions about their treatment plan, and supports the model put forward by Walsh KE et al. 2011 that describes parents and caregivers as a final layer of

protection for children from medication errors. However, there is a paucity of literature on the role of parents in preventing or contributing to medical errors (Walsh KE et al. 2011).

Avery AJ et al. 2013 report that many inter-related contributory factors underpin prescribing errors in UK general practice including: prescriber training, knowledge and experience of specific medications and patients, risk perception, patient characteristics, team collaboration, GPs signing nurse generated prescriptions without seeing the patient, poor working conditions including time pressures and interruptions, and computer-related issues. Several of these factors were also present in the student's study. Poor working conditions were an important reported factor, not only in general practice but also in community pharmacies, other important reported factors include inadequate staff knowledge about certain medications and patient characteristics such as age.

Staff failure to follow protocol was common described as contributing to medication incidents, and although failure to follow National Institute of Health and Care Excellence or Scottish Intercollegiate Guidelines Network (SIGN) epilepsy treatment protocols was uncommon in the national epilepsy review, compliance was noted as sub-optimal.

5.2.3 Recommendations for improvement

The electronic prescription service (a form of CPOE) was rolled out in UK General Practice in 2005 and is now widely used. Introduction of such technology may have reduced certain types of medical error, as it did in the paediatric in-patient setting, but its impact on safety must be regularly evaluated, as it may predispose to IT-related medical errors at the human-machine interface (Conroy S et al. 2007; Walsh KE et al. 2005; Wong IC et al. 2009). The electronic prescription service could be improved with better linkage to patient records, enabling more sensitive detection of medication contraindications specific to each patient, and by incorporating more rigid forcing functions into it e.g. to stop prescribers ignoring or overriding certain high-risk safety alerts (Kaushal R et al. 2001).

Although prescribing errors often preceded reported dispensing errors, errors at the prescribing-dispensing interface were more frequent i.e. the prescription was correct but a dispensing error occurred. Electronic transmission of prescriptions from the prescriber to the dispensing community pharmacy has been proposed to address errors at the prescribing-dispensing interface. However, if not implemented correctly medication errors could increase, as highlighted by a recent review of this technology in UK primary care (Franklin BD et al. 2014).

Education and training of all pharmacy staff, in human factors for example, would facilitate staff to recognise error prone-areas of their practice, and to strengthen those areas (Campino A et al. 2009; Kaji AH et al. 2006; Leonard MS et al. 2006; Levine SR et al. 2001; Sturgess E et al. 2011; Sullivan MM et al. 2010). Providing staff with the tools to address safety problems themselves is a more effective and flexible way to address problems that may be specific to their pharmacy.

Implementation of a barcoding system for all dispensed medication could reduce the potential for human error, by acting as an additional safety check prior to giving the patient their medication (Kaushal R et al. 2001; Morriss FH et al. 2009; Poon EG et al. 2010).

Publicising community pharmacy error rates could also facilitate medication error reduction, as transparency incentivises improvement (Sturgess E et al. 2011). This form of feedback could encourage competition between pharmacies and help to create a culture of openness and shared learning where the more error-prone pharmacies learn from the 'safer' pharmacies (Eisenhut M et al. 2011;Leonard MS et al. 2006).

Children with chronic conditions requiring regular treatment, such as those with epilepsy, should have written management plans that are accessible to all relevant parties. The RCPCH have since launched an epilepsy passport to improve clarity about epilepsy management, including any changes to the treatment plan, to improve continuity of care for children seeing multiple professionals.

5.3 Discussion of diagnosis-, assessment-, referral-, and communication-related findings

5.3.1 Summary

A considerable proportion of diagnosis and assessment-, referral-, and communication-related incidents were associated with telephone assessments, and underpinned by similar factors requiring attention. Therefore, in this section, these incidents will be considered together in the context of current literature and recommendations for improvement.

Diagnosis, assessment, referral, and communication incidents were most commonly reported in younger children (less than three years old), presenting acutely with injuries, non-specific signs and symptoms such as fever and altered consciousness, and skin conditions such as rashes and discolouration.

These 'unwell' children frequently experienced inadequate telephone triaging, and delays in assessment and referral. Incidents underpinning diagnosis, assessment and referral failures included: communication incidents and other assessment failures, particularly inadequate history taking. Communication incidents (primary incidents and those contributing to diagnosis, assessment and subsequent referral incidents) were related to inadequate safety netting, and provision of the wrong advice to caregivers during telephone assessment.

Caregivers were described as playing an important role in incidents related to telephone assessment. They contributed to incidents through poor interpretation of their child's conditions, and subsequent communication of that inappropriate interpretation to the telephone assessor. However they also prevented incidents by challenging the management decisions of staff. Other factors contributing to telephone assessment failures were related to issues with protocolised medicine, including failing to follow protocols such as those related to safety netting, failures of the protocols themselves, and poor staff critical thinking whilst using the protocols.

5.3.2 Context of current literature

5.3.2.1 Telephone assessment

The finding that younger children were more prone to reported failures involving diagnosis, assessment, referral, and communication is not unexpected and likely reflects: differences in physiological reserve and speed of deterioration, differences in disease epidemiology, communication difficulties, and dependency of younger children on caregivers to recognise illness (Walsh KE et al. 2014;Wolfe I et al. 2011). In addition, considering the high proportion of incidents involving telephone assessment and that those under-5 years old are the highest users of NHS direct (contributing to 25% of all calls), it is unsurprising that most diagnosis, assessment, referral, and communication incidents in this study involved younger children (Cook EJ et al. 2013).

Problems with telephone assessment (which occasionally resulted in referral incidents) were prevalent in this dataset, and concerns about the safety of telephone triaging are echoed in the literature (Derkx HP et al. 2008; Giesen P et al. 2007; Huibers L et al. 2011; McLellan N 1999; McLellan N 2004; O'Cathain A et al. 2003; Smits M et al. 2010; Stewart B et al. 2006). Given the focus of this study, 'unwell' children in primary care, and considering that NHS direct handles over 500,000 calls a month with up to 40% involving children, numerous reports of telephone assessment problems were anticipated (McLellan N 2004; Stewart B et al. 2006).

Safe telephone assessment involves correctly determining the urgency of the child's condition, and subsequently giving appropriate management advice, which may involve referral to emergency services. A systematic review of telephone triaging conducted by Huibers L et al. 2011, highlights that approximately 10% of calls are unsafe, and that patients presenting with highly urgent symptoms may be particularly vulnerable to unsafe care. Triaging incidents in the student's study included both under and overestimation of urgency. Derkx HP et al. 2008 highlight similar issues with telephone triaging, and that in 41% of calls urgency is underestimated, and that in 1% urgency is overestimated. Overestimating the urgency of a child's condition is also a safety concern as it may result in unnecessary referrals and further investigations (Cook R et al. 2010).

The high number of incidents involving young children with injuries, non-specific signs and symptoms, and skin conditions described in this dataset, is in keeping with the literature (Cook EJ et al. 2015;Cook EJ et al. 2013). Cook EJ et al. 2013 highlight that most NHS direct calls involving children are for: crying (1.01 calls/100 population per annum), skin hand or nail conditions (0.9 calls/100 population per annum), and wounds and injuries (0.67 calls/100 population per annum).

Paediatricians have expressed concern about the use of telephone assessment for children due to the non-specific nature of many childhood illnesses, in addition to dependency on caregivers to observe their child and most importantly to interpret their observations and communicate those interpretations effectively (McLellan N 1999;McLellan N 2004;Stewart B et al. 2006). The student's study supports these concerns, because misunderstandings between caregivers and professionals over the telephone, such as misunderstanding fever management advice or the difference between a blanching and non-blanching rash, were frequent.

Many highlight issues with the CDS used by NHS direct, particularly in relation to assessing the urgency of children's conditions (McLellan N 1999;McLellan N 2004;Stewart B et al. 2006). CDS software is designed to minimise risk by improving the consistency of assessment and decision making during telephone triaging, but they also reduce professional autonomy, a factor underlying many incidents in this study (McLellan N 2004;O'Cathain A et al. 2003). However, the trade-off between their sensitivity to urgent presentations requiring emergency management, and their specificity (to exclude non urgent cases from referral to emergency services), and their responsiveness to contextual information, is contentious and dependant on the CDS software used. Many argue that it is unclear whether CDS can compensate for inadequate knowledge and clinical paediatric experience (Doctor K et al. 2014;Leprohon J and Patel VL 1995;McLellan N 1999;McLellan N 2004;Monaghan R et al. 2003;O'Cathain A et al. 2003).

Staff are able to submit suggested improvements to the CDS software and calls are regularly audited against minimum quality standards. However, without evaluating, on a large scale, the outcomes of children triaged via this process, it is unclear whether auditors can reliably detect inadequacies in the CDS software.

Huibers L et al. 2011 highlight the importance of good quality history taking to inform triaging decisions, a factor which commonly contributed to triaging incidents in this study. Healthcare professionals' ability to assess and manage 'unwell' children is influenced by their experience and professional background. Monaghan R et al. 2003 highlight differences in triaging practices between general nurses and children's nurses (Pettinari CJ and Jessopp L 2001;Smith S 2010). Despite this, only 1% of professionals employed by NHS direct have a background in paediatrics (McLellan N 2004). It is therefore unsurprising that in relation to paediatric triaging, nurses are reluctant to deviate from and challenge the triaging protocols and algorithms. This was reflected by the high number of reports describing poor staff critical thinking as contributing to diagnosis, assessment, and referral incidents. Smith S 2010 also reiterates that clinical reasoning (i.e. critical thinking and adherence to guidelines and protocols) frequently underpins safety failures related to triaging, diagnosis and treatment in primary care out-of-hours.

5.3.2.2 Safety netting

Communication incidents involving caregivers were frequently related to: misunderstandings between caregivers (callers) and professionals, the use of inappropriate triaging protocols, or not following protocols or safety netting guidelines. Derkx HP et al. 2008 also report problems with misinterpretation of callers' responses to triaging questions leading to assessment incidents. Despite communication incidents being highlighted as a priority area requiring improvement in primary care, there is a paucity of literature on the topic, particularly in relation to telephone triaging-related communication errors involving caregivers (Cresswell KM et al. 2013; Rees P et al. 2015). However, NHS direct safety netting has been described as generic and not specific to the child or family (Roland D et al. 2014).

5.3.2.3 Role of caregivers

The role of patients and caregivers in preventing safety incidents, although not a new concept in healthcare quality and safety, has been relatively underexplored in the literature (Berger Z et al. 2013; Walsh KE et al. 2005). Roland D et al. 2014 theorise that, providing comprehensive safety netting advice, that includes

information on diagnostic probabilities and uncertainties, could empower patients and caregivers to challenge the diagnostic process as they feel is necessary. They believe that educating patients in this manner will improve their understanding of the process, and additionally encourage them to provide pertinent information, for example about the history of their presentation, which may otherwise have been overlooked. Educating caregivers in this manner is supported by a randomised controlled trial, which reports improved identification of serious symptoms in educated patients (McCarthy PL et al. 1990).

Parents are often disempowered by healthcare, despite having the potential to be a valuable asset in their child's care(Graedon J and Graedon T 2006; Neill SJ 2010; Titcombe J 2015). Parents are a reasonably constant presence in their child's life, they know their child's normal behaviour and temperament, and are particularly astute observers when their child is unwell (Brady PW et al. 2014; Graedon J and Graedon T 2006). Titcombe J 2015 argues that parents will "always have continuity and context on their side", we must therefore not overlook their opinion. They are important allies. The role of caregivers is being increasingly valued in healthcare: many paediatric early warning scores take into account parental concern as an indicator of the severity of a child's illness, and Birmingham Children's Hospital permit parents to write in the child's medical notes (Roland D 2015; Roland D et al. 2013; Sen AI et al. 2013; Titcombe J 2015).

Brady PW et al. 2014 demonstrate the value of involving parents in the hospital care process by allowing them to directly activate the rapid response team. They report that families use the system responsibly, there are few unnecessary activations, 24% of activations result in intensive care transfers, and fewer cardiorespiratory arrests occur.

5.3.4 Recommendations to improve telephone assessment

5.3.4.1 Education and training

Mandatory paediatric training for all trainee GPs, to increase their tacit knowledge through experiential learning, is imperative to improve the detection of seriously unwell children presenting in person or via the telephone to general practice (Rees P et al. 2015; Wolfe I et al. 2011). All healthcare professionals in contact with children should be familiar with the 'spotting the sick child' website, and complete the online modules as part of their continuing professional development. NHS direct (now NHS 111) calls are regularly audited and the results of that audit are fed back to staff, peer-review rather than senior-review of calls could help reviewers and those being reviewed to maintain high standards and prevent them from developing poor habits over time (Graber ML et al. 2012; Singh H et al. 2010; Thammasitboon S and Cutrer WB 2013).

All healthcare professionals, whether GPs or telephone assessment staff, should be made aware of error-prone areas of practice, such as triaging the wrong symptom in a child with multiple symptoms over the telephone (Graber ML et al. 2012; Singh H et al. 2010; Thammasitboon S and Cutrer WB 2013).

5.3.4.2 Teamwork/ collaboration

This study supports re-organisation of general practice for children into child health hubs, which is currently being piloted in some parts of the UK (Wolfe I et al. 2011). This involves co-locating paediatricians, GPs with an interest in paediatrics, and other members of the multi-disciplinary team, enabling integrated care and increasing the support and expertise readily available to professionals caring for children in the community, in and out of hours.

5.3.4.3 Amend protocols

Telephone assessment protocols should be regularly reviewed and updated. It is insufficient to depend on professionals to detect and report inadequacies in the CDS software, particularly when they are unaware of patient outcomes. The outcomes of children assessed via the telephone should be regularly reviewed

(rather than simply auditing the calls themselves) to assess the adequacy of protocols. The sensitivity of protocols for certain groups of patients could then be improved through amending the CDS software (Graber ML et al. 2012;Ramnarayan P et al. 2006a;Ramnarayan P et al. 2004;Ramnarayan P et al. 2006b;Singh H et al. 2012). Regular reviews of calls to measure error rates would allow feedback to professionals, but also, when analysed in conjunction with outcome data, would enable assessment of whether changes such as updates in CDS protocols actually improve patient safety (Ramnarayan P et al. 2006a;Ramnarayan P et al. 2004;Ramnarayan P et al. 2006b;Singh H et al. 2012).

Certain error-prone presentations may benefit from mandatory senior assessment. Also CDS could be strengthened to improve error-prone areas of practice, for example by inserting reminders when triaging head wounds to double check the absence of a head injury (which would require triaging with a different protocol) (Graber ML et al. 2012;Ramnarayan P et al. 2006a;Ramnarayan P et al. 2004;Ramnarayan P et al. 2006b;Singh H et al. 2012).

5.3.4.4 Patient/ caregiver empowerment

The utility of caregivers must not be under-estimated, they should be directed to evidence-based resources to get further information about their child's condition and when it is and is not appropriate to present to healthcare professionals (Singh H et al. 2010).

5.3.4.5 Better safety netting

This study points to a clear need for improved safety netting via the telephone. Parents and caregivers should receive oral and written safety netting information (perhaps via e-mail, text messages, or smart phone applications) (Roland D et al. 2014). The content of safety net advice must be reviewed in this setting, to ensure it is individually tailored and comprehensive to include: diagnostic probabilities, warning signs to be aware of, a likely time frame for illness recovery, and advice on the most appropriate way to access healthcare if there is no improvement or the child deteriorates (O'Cathain A et al. 2003). Safety netting protocol adherence could be improved through using mnemonics or checklists (Kim SW et al. 2012;Sahyoun C et al. 2013;Starmer AJ et al. 2013;Weingart C et al. 2013a).

5.4 Recommendations for future research

This study has highlighted priority areas requiring improvement using incident report data, however estimates of the burden of unsafe primary care for children would be beneficial. Mangione-Smith R et al. 2007 in the USA were the first to measure ambulatory care quality for children; this study is now being repeated in Australia (Hibbert PD et al. 2015). The UK would also benefit from a comprehensive and generalisable case note review, to determine which areas of primary care are most harmful and costly to child health, in order to target improvement interventions efficiently to those areas most in need of improvement (Hibbert PD et al. 2015;Mangione-Smith R et al. 2007a).

With the emergence of large datasets such as those managed by the Health and Social Care Information Centre (HSCIC) and the Secure Anonymised Information Linkage (SAIL) databank, that can be linked to provide epidemiological data on care quality, there is considerable scope for future studies to evaluate the epidemiology of healthcare harm in children (Dattani N et al. 2013; Hardelid P et al. 2013; Hardelid P and Gilbert R 2013). Future research could utilise such resources to model the risk of substandard care given certain characteristics such as age, location, past medical history, treatment history, family history, and social factors including family socioeconomic status.

The role of caregivers in contributing to, and mitigating, substandard care was a prominent theme in this dataset. Future work should consider the benefit of parental involvement in quality improvement projects. The role of caregivers as potential 'error safety nets' should be explored, particularly in the context of them having access to their children's medical records, which is anticipated imminently (Woodman J et al. 2015).

Finally, this study hypothesises that socially vulnerable children are disproportionately vulnerable to unsafe and poor quality care. This hypothesis should be investigated in future research, to include an in-depth exploration of the reasons for such a potential pre-disposition, and piloting interventions targeted to improve the safety of care for this vulnerable population of children, by addressing contributory factors specific to this group.

Chapter 6: Conclusions

This thesis explores the quality and safety of primary care for children by identifying key reported safety issues related to childhood vaccination and the provision of care to 'unwell' children.

Priority areas related to vaccination requiring improvement include administration of: the wrong number of doses, vaccines at the wrong time, and the wrong vaccine. Weaknesses underlying these incidents include documentation failures, appointment management problems, staff mistakes, and failure to follow vaccination protocols. Certain groups of children such as looked-after children appeared disproportionately vulnerable to these failures. Recommendations to potentially address these issues include: building IT infrastructure to mitigate vaccination-related documentation weaknesses such as implementing electronic 'red' books; and encouraging better communication and shared decision making between healthcare professionals and caregivers.

Key primary care-related safety incidents involving 'unwell' children were related to medication provision; and diagnosis, assessment, referral, and communication failures with regards to telephone assessment.

Medication-related incidents were underpinned by: staff factors such as mistakes and failing to follow protocols; organisational factors such as poor working conditions and inadequate care plans and protocols; patient factors such as age; and medication factors such as adjacent storage of similar medications. Recommendations to mitigate these incidents include: improving linkage between electronic prescription services and patient records, routine electronic transmission of prescriptions to dispensing pharmacies, barcoding of all medications to act as a double-check prior to dispensing, and providing human factors training to all pharmacy staff.

Telephone assessment failures were the result of assessors choosing the wrong assessment protocols, not utilising the protocols correctly, or not using critical thinking to challenge inappropriate protocol outcomes. Inadequate safety netting was also described in this context. Given the nature of these incidents, parents and

caregivers played an important role in ensuring their children received appropriate telephone assessments.

Recommendations for improving telephone assessment of 'unwell' children include: reviewing and amending CDS software and telephone assessment protocols, to ensure that they are comprehensive, sensitive, user-friendly, and suitable for use on children. The quality of safety netting protocols should also be reviewed and adherence to them should be improved, perhaps through the use of a mnemonic.

Further studies are required to assess the burden of unsafe primary care in children; and to test the hypotheses generated from this study with regards to the nature of unsafe primary care, the causative factors underlying this unsafe care, and the recommendations for improvement.

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Appendix 1 - Published manuscripts

1.1 BMJ letter (contraindicated BCG vaccination)

Rees P, Evans H, Panesar S, Llewelyn M, Edwards A, Carson-Stevens A. Contraindicated BCG vaccination in "at risk" infants. BMJ. 2014 Sep 9;349:g5388. doi: 10.1136/bmj.g5388





BMJ 2014;349:g5388 doi: 10.1136/bmj.g5388 (Published 9 September 2014)

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LETTERS

BCG VACCINATION AND TB IN CHILDREN

Contraindicated BCG vaccination in "at risk" infants

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Roy and colleagues highlight the effectiveness of the BCG vaccine in preventing tuberculosis in immunocompetent children.1 However, those responsible for prescribing and administering BCG vaccinations must be mindful of the contraindications for immunocompromised children.²

Our study of primary care related paediatric safety incidents from the England and Wales National Reporting and Learning System identified 1790 vaccine related incidents during 2005-13.3 Despite under-reporting of safety incidents, we found 99 reports of BCG related incidents in children at risk of tuberculosis exposure—by living in or having family members from areas with a high prevalence of tuberculosis. A recurrent theme was administration of BCG vaccines to infants of HIV positive mothers in whom full assessment of their HIV status had not been completed. Children infected with HIV are at risk of disseminated BCG disease.² Public Health England advises that BCG vaccinations are given only after two appropriately timed negative HIV tests.4 Poor communication between health services—maternity care, child health, and general practice—is often implicated. Typically, infants are sent appointments by child health services as a result of deficiencies or discrepancies between maternal and child health records. This creates a situation whereby healthcare professionals are unaware of a child's vulnerability.

Vaccine errors remain a threat to patient care.5 Contraindications to BCG must be explicitly stated within parent held records. Better linkage of maternal and child health records that are accessible in primary and secondary care is also needed.

Healthcare professionals must not rely solely on medical records but should consider, double check, and fully explore with parents (including use of translation services if needed) the child's suitability for the vaccine before administration.

Competing interests: AC-S and AE are co-chief investigators of a National Institute for Health Research Health Services and Delivery Research grant to characterise patient safety incident reports in primary care. PR is a research assistant employed to work on the study. HE and ML have no conflicts of interest. SP is a former clinical adviser at the National Patient Safety Agency (2008-10) and an academic clinical fellow at Imperial College London working for the National Reporting and Learning System research programme.

Full response at: www.bmj.com/content/349/bmj.g4643/rr/762808.

- Roy A, Eisenhut M, Harris RJ, Rodrigues LC, Sridhar S, Habermann S, et al. Effect of BCG vaccination against Mycobacterium tuberculosis infection in children: systematic review and meta-analysis. BMJ 2014;349:g4643. (5 August.)
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- National Institute for Health Research. Characterising the nature of primary care patient safety incident reports in England and Wales: mixed methods study. www.nets.nihr.ac. uk/projects/hsdr/1264118.
 Public Health England. Contraindications and special considerations. In: Immunisation
- against infectious disease. 2013:41-8. www.gov.uk/government/publications/ contraindications-and-special-considerations-the-green-book-chapter-6. Derrough TF, Kitchin NR. Occurrence of adverse events following inadvertent
- ration of childhood vaccines. Vaccine 2002;21:53-9.

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1.2 Archives of disease in childhood letter (socially deprived children)

Rees P, Edwards A, Powell C, Evans H, Panesar S, Carson-Stevens A. Disparities in the quality of primary healthcare for socially deprived children. Arch Dis Child. 2015 Mar;100(3):299-300. doi: 10.1136/archdischild-2014-307618.

LETTERS

Disparities in the quality of primary healthcare for socially deprived children

Mbeledogu et al1 demonstrate a reduction in unintentional poisonings in children in England, yet highlight the protracted social disparities among these children. This resonates with our exploration of over 20 000 paediatric safety incidents from primary care in England and Wales where the 'Inverse Care Law' features prominently.² These reports include: looked-after children, immigrants, travellers, parents with addiction problems and those belonging to other disadvantaged groups. Data mining identified over 500 reports involving these groups and 33 reports involving homeless children or families. Such a reporting system has not previously been used to assess unsafe care in socially deprived groups.

Thematic analysis of free-text in reports involving homeless children was undertaken and five overarching failures were identified (see table 1). Some disclosure failures resulted in breaches of confidentiality where the location of an abused child was revealed to their abusers through administrative errors (see table 1, example 1). Conversely, uncertainty around when to disclose information to members from other disciplines or services was also reported (see table 1, example 2). This led to service fragmentation (eg, no health visitor or general practitioner input) and poor continuity of care

(eg, missed safeguarding visits) for at-risk families. Difficulties accessing primary and social care featured often, where some general practices refused to register children with temporary addresses—a typical problem for the homeless (see table 1, example 3). In addition, the child was unable to access care if their mother was not concurrently registered at the practice. Example 4 illustrates how this culminated in a child not receiving primary healthcare for 4 months. Frequent relocation of families resulted in untimely transfer of records. In example 5, an at-risk child was without health visitor support for several weeks.

Our thematic analysis of 33 reports has identified several important issues for consideration by healthcare professionals and policymakers. We recommend general practices review their policies on child registration and have a process in place to make exceptions to any rules, for example requiring co-registration with a parent, for vulnerable children. In addition, all healthcare professionals must be aware of what constitutes a confidentiality breach. Finally, alerting an abuse perpetrator to the location of a victim of abuse should be considered a 'never event'. Never events are unacceptable and preventable incidents that could result in severe harm or death. with defined criteria, and must be reported to the Department of Health.

It is unclear whether these are isolated incidents disclosed within the free-text, or systemic failures particular to homeless families since important patient characteristics (eg, socioeconomic status) are missing from most reports. The value of data linkage to evaluate the impact of patient characteristics on healthcare outcomes was demonstrated in a recent UK-wide enquiry into child mortality.4 From our characterisation of reports involving children, insights for prioritising and designing future safety interventions could be gained by linking incident reporting systems with electronic medical records and other public or social care registries. This would enable the identification of incident reports relevant to specific groups.

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Competing interests AC-S and AE are co-chief investigators of a NIHR HS&DR grant to characterise patient safety incident reports in primary care. PR is a research assistant employed to work on the study. HE and CP have no conflicts of interest. SP is a former clinical adviser at the NPSA (2008-2010) and an Academic Clinical Fellow at Imperial College London working for the NRLS research programme.

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Table 1 Opportunities for health and social care improvement and free text examples

Free-text extracts Opportunity for improvement Breaches of confidentiality Example 1. 17 year-old victim of domestic abuse had been an inpatient and is pregnant. During admission a referral was sent to ultrasound. This patient was discharged to a refuge for safety, and her address changed I later noticed that her address was changed back-to that of the perpetrator of the domestic abuse-by staff in radiology. This patient's appointment has been sent to the perpetrator's address. Health and social care professional uncertainties around Example 2. A health visitor attempted a home visit for a baby on the child protection register for neglect. when to disclose information Information regarding the mother and baby being moved to a refuge were withheld. The social worker was reluctant to give the forwarding address and contact number to the health visitor. Example 3. A Social Services department refused to accept a referral as the refugee is only at a temporary address. The family intend to stay in *** this was communicated to the *** department who still declined to Inability to access or register for primary care using a temporary address accept the referral. Inability to access primary care if mother is not Example 4. The patients' mother fled domestic violence, registered as homeless and was stabilised on methadone. Registration was arranged with Surgery A, who have subsequently de-registered the child as mother concurrently registered requires on-going support for her addiction [not available at that practice]. Centre B stated that the child was outside of their catchment area. Centre C cannot register a child without the parent (they cannot offer opiate substitution). Therefore a child in contact with services for 4 months is not registered for primary care. This child known to social services does not have a named social worker. Example 5. A family were rehoused in the local area with no indication of vulnerability. On examination of the Untimely transfer of records on relocation notes, I found that the family was known to social services with concerns regarding the safety of the child. The support team was contacted but the family were not known to them. The family moved from *** to ***. Note: support team was contacted but the family were not known to them. The family moved from were sent to a health visitor for the homeless but the family had already moved again. Therefore a vulnerable family with a young baby were without support for weeks, putting a child at risk.

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1.3 Lancet letter (UK child mortality)

Rees P, Panesar SS, Edwards A, Carson-Stevens A. Child mortality in the UK. Lancet. 2014 Nov 29;384(9958):1923-4. doi: 10.1016/S0140-6736(14)62272-8.

Child mortality in the UK

The UK did particularly poorly in terms of the mortality estimates for children younger than 5 years, produced by the Global Burden of Diseases team (Sept 13, p 957). At 4-9 deaths per 1000 in 2013, the UK has almost the highest rate in western Europe, double that of Sweden, a country with one of the lowest mortality estimates. Furthermore, high-income countries with a high proportion of children living in relative poverty, have a high rate of under-5 mortality (figure). 12

Child poverty in high-income countries varies greatly, suggesting that the level of poverty might be a policy choice, amenable to change by national governments. Analyses consistently show that much of this variation is directly related to differences in tax and benefit systems.³ Unfortunately, signs of concern suggest that present policies in the UK might be making the situation worse.⁴

Absolute child poverty increased for the first time in 17 years in the UK, during changes to the tax and benefit system that are reducing the adequacy, eligibility, and access to benefits, especially for some families with lowest incomes with children.⁴ Governmental cuts to spending for the public sector are affecting services relied on by families with low incomes who have children, with the largest spending cuts to budgets of local authorities in deprived areas.⁵

If we are to reduce the number of child deaths in the UK we need a welfare system that prioritises children, with a renewed focus on improving the circumstances in which children grow up, alongside systematic improvements in health services.⁶

We declare no competing interests

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Wang H, Liddell CA, Coates MM, et al. Global, regional, and national levels of neonatal, infant, and under-5 mortality during 1990–2013: a systematic analysis for the Global Burden of Disease Study 2013. Lancet 2014; 384: 957–79.

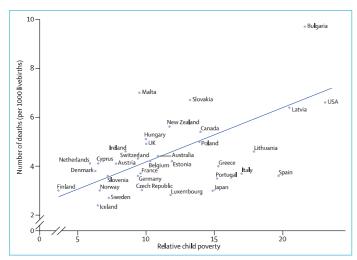
- 2 UNICEF. Child well-being in rich countries: a comparative overview. 2011. http://www. unicef.org.uk/Images/Campaigns/FINAL_RC11-ENG-LORES-fnl2.pdf (accessed May 10, 2014).
- 3 Marx I, Nelson K. Minimum income protection in flux. Houndmills, Basingstoke, Hampshire, UK: Palgrave Macmillan, 2013.
- 4 Taylor-Robinson D, Harrison D, Whitehead M, Barr B. Doctors need to take the lead on poverty's effects on health. BMJ 2013; 347: 7540.
- 5 Taylor-Robinson D, Gosling R. English north-south divide. Local authority budget cuts and health inequalities. BMJ 2011; 342: d1487.
- 6 Wolfe I, Thompson M, Gill P, et al. Health services for children in western Europe. Lancet 2013; 381: 1224–34.

Haidong Wang and colleagues1 emphasise the importance of global monitoring to identify countries most in need of improvement. Although we support this ambition, we recognise that every country has a responsibility to build infrastructure and assimilate data sources (old and new) to identify opportunities to improve the quality of health and provision of social care for children. A review² suggested that more than 20% of child deaths in the UK in 2013 had so-called modifiable factors, whereby health care could have intervened to mitigate the risk of death. Analysis of routine data sources, such as data from incident reports for patient safety, could help to understand what common contributory, modifiable factors might be.3

A global registry for paediatric safety using a minimum dataset from every country, irrespective of economic setting, would allow the necessary surveillance to ensure harm was mitigated against and indeed solutions developed to prevent these avoidable deaths in children. This registry will need a combination of analytics with clinical expertise to generate action-orientated outputs with strong face validity in the health-care profession.

AE, AC-S, SSP, and PR work on a project funded by the National Institute of Health Research (NIHR) Health Services and Delivery Research (HS&DR) to characterise patient safety incident reports in primary care. The views and opinions expressed therein are those of the authors and do not necessarily reflect those of the HS&DR, NIHR, UK National Health Service, or the Department of Health (UK).

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 $\emph{Figure}: Child poverty and mortality in 35 countries in the Organisation for Economic Co-operation and Development}$

Child relative poverty rates (having equivalised household income of less than 50% of the national median) are from the 2013 UNICEF report. Mortality estimates are for children younger than 5 years for 2013 from the Global Burden of Diseases analysis by Wang and colleagues. Linear regression trend line; mortality in children younger than 5 years is calculated by $1.94 + 0.22 \times \text{relative}$ child poverty; p < 0.0001; R2 = 0.40.

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1.4 Pediatrics manuscript (safety incidents in general practice)

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Safety Incidents in the Primary Care Office Setting

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BACKGROUND: In the United Kingdom, 26% of child deaths have identifiable failures in care. Although children account for 40% of family physicians' workload, little is known about the safety of care in the community setting. Using data from a national patient safety incident reporting system, this study aimed to characterize the pediatric safety incidents occurring in family practice.

METHODS: We undertook a retrospective, cross-sectional, mixed methods study of pediatric reports submitted to the UK National Reporting and Learning System from family practice. Analysis involved detailed data coding using multiaxial frameworks, descriptive statistical analysis, and thematic analysis of a special-case sample of reports. Using frequency distributions and cross-tabulations, the relationships between incident types and contributory factors were explored.

RESULTS: Of 1788 reports identified, 763 (42.7%) described harm to children. Three crosscutting priority areas were identified: medication management, assessment and referral, and treatment. The 4 incident types associated with the most harmful outcomes are errors associated with diagnosis and assessment, delivery of treatment and procedures, referrals, and medication provision. Poor referral and treatment decisions in severely unwell or vulnerable children, along with delayed diagnosis and insufficient assessment of such children, featured prominently in incidents resulting in severe harm or death.

CONCLUSION: This is the first analysis of nationally collected, family practice—related pediatric safety incident reports. Recommendations to mitigate harm in these priority areas include mandatory pediatric training for all family physicians; use of electronic tools to support diagnosis, management, and referral decision-making; and use of technological adjuncts such as barcode scanning to reduce medication errors.

abstract



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WHAT IS KNOWN ON THIS SUBJECT: More than a quarter of child deaths in the United Kingdom are estimated to have identifiable failures in care. Although children account for 40% of the family practice workload, little is known about iatrogenic harm to children in this setting.

WHAT THIS STUDY ADDS: This is the first analysis of nationally collected pediatric safety incident reports from family practice. To mitigate harm to children, priority areas requiring improvement include medication provision, referral of unwell children, provision of evidence-based treatment, and adequate diagnosis and assessment.

ARTICLE

The quality and safety of health care is an established public health concern.1-3 Efforts to improve health care delivery have largely been in high-income countries and in hospital rather than community settings.4 There has also been relatively little focus on child health and pediatric services.5 Although children account for 40% of family physicians' workload, little is known about the safety of care delivered to children in the community setting.6,7 In the United Kingdom, family practice is usually the patient's first point of contact with the health care service; most health care encounters occur in this setting, and it acts as a gateway to acute and specialist hospital-based services.

Several methods have been used to measure health care-associated harm in pediatrics, and challenges remain.8 A child mortality review in the United Kingdom found that 43% of child deaths had potentially avoidable factors, and 26% had identifiable failures in care. 9,10 However, because of the relatively low number of deaths, mortality data are not powered to provide epidemiologic associations on the impact of patient safety issues.11 In contrast, incident reporting systems have a well-established infrastructure, can provide large quantities of data on pediatric safety, and have the potential for greater insight and hypothesis generation than mortality data alone.12

This study is the first systematic, mixed-methods analysis of nationally collected patient safety incident reports involving children in family practice. Previous studies have demonstrated the value of using incident reporting systems to improve care quality in neonatal settings.13 National repositories of patient safety incident reports permit the identification of clusters of similar reports that could be potentially overlooked as rare events at a local level. Priority areas for practice can be identified at a national level, and insights can be gained from exploring

common contributory factors related to incident type. These can inform the basis of recommendations for improvements in pediatric care.^{5,14,15}

This study aimed to explore the nature and severity of pediatric safety incidents occurring in family practice and their potential contributory factors. Our objective was to identify potential priority areas for intervention.

METHODS

National Reporting and Learning System

The England and Wales National Reporting and Learning System (NRLS) was established in 2003 as a national repository of reports about patient safety incidents, defined as "any unexpected or unintended incident[s] which could have, or did, lead to harm to one or more patients."16 The NRLS receives ~100 000 reports per month written by patients or health care providers from any National Health Service organization in England and Wales, and annually >65 000 reports involve children.5 Reports can be submitted directly to the NRLS online, or staff can report incidents to their local health boards, which subsequently upload the reports to the NRLS; all reports are anonymized. Each NRLS report contains several categories of data, including patient age, incident location, date of incident, free-text fields containing

descriptions of the incident, and severity of harm outcome (no harm, low harm, moderate harm, severe harm, and death).

Sample selection

Incident reports from family practice in England and Wales (April 2003 to June 2012) involving patients aged <18 years were identified by manually applying an age filter to all 46 902 family practice reports available. On reading the reports, some were still evident as adult cases and excluded.

Methodology

We undertook a retrospective, crosssectional, mixed-methods study combining a detailed data coding process, frequent generation of data summaries using descriptive statistical analysis, and thematic analysis of a theoretical and specialcase sample of reports. New ideas and hypotheses emerged throughout each step of analysis for later corroboration.

Data Coding

We used an inductive, grounded approach to apply codes to each incident report from a codebook containing 2 distinct multiaxial coding frameworks (empirically developed in-house) to describe the type of safety incident (administration, medication, etc.) and contributory factors (patient, staff, environmental, etc.), as well as the World Health Organization (WHO)

TABLE 1 Severity of Harm Described Within Reports

Severity of Harm	Reports, n (%)
No harm: patient outcome is not symptomatic and no treatment is required	1025 (57.3)
Low harm: patient outcome is symptomatic, symptoms are mild, loss of function or harm is minimal and intermediate but short term, and no or minimal intervention is required	675 (37.8)
Moderate harm: patient outcome is symptomatic requiring intervention, an increased length of stay, or causing permanent or long-term harm or loss of function	71(4)
Severe harm: patient outcome is symptomatic, requiring life-saving intervention or major surgical/medical intervention, shortening life expectancy or causing major permanent or long-term harm or loss of function	9 (0.5)
Death: on balance of probabilities, death was caused or brought forward in the short term by the incident	8 (0.4)
Total	1788 (100)

International Classification of Patient Safety to describe harm severity (see Table 1 for definitions).^{17,18}

We concluded from early pilot work that existing classification systems for primary care did not permit us to code at the level of detail needed to answer our research questions. Thus, we empirically developed 2 coding frameworks to describe the incidents in detail and capture the complex and often multiple contributory factors. We acknowledge that at initial conceptual level, preexisting taxonomies (largely the WHO International Classification of Patient Safety and the Learning from International Networks about Errors and Understanding Safety in Primary Care [LINNAEUS Euro-PC] taxonomy) informed our high-level considerations for both frameworks, as we recognized this was important to permit comparison between studies of this nature.17,19

Initially, an iterative constant comparative method was used: a small number of incident narratives were read, preliminary codes were extracted, and these codes were revised through further reading of additional reports. This cycle was repeated until a stable set of codes was developed between 2 authors (Miss Rees and Dr Carson-Stevens, both trained in root cause analysis and human factors). Subsequent codes were discussed and frameworks were iterated at weekly coding meetings involving family physicians, health services researchers, and anthropologists.20 This process enabled development of codes particular to reports about children in family practice.

One to 4 codes were used to describe the incidents and 1 to 4 codes to describe the potential contributory factors, in keeping with 9 recursive incident analysis rules developed by the Australian Patient Safety Foundation (see Supplemental Appendix 1).²¹ Twenty percent of the reports were independently double-

coded (Miss Rees and Dr Carson-Stevens). Interrater reliability (Cohen's κ statistics) were calculated for the primary incident type (chronologically closest to the outcome experienced by the patient) and contributory factors. These were $\kappa = 0.88$ (95% confidence interval 0.85-0.9) and $\kappa = 0.86$ (95%) confidence interval 0.82-0.9), respectively. Disagreements in coding were resolved by a third reviewer (Dr Panesar) and discussion with the coding team. Coding of free-text entries allowed categorization of reports by incident type to provide the basis for subsequent data analysis. Similar methods have been used in previous research involving hospital NRLS reports.22,23

Where the severity of harm stated in the NRLS report conflicted with the report's free-text incident description, the level of harm was adjusted using the WHO's International Classification of Patient Safety definitions.¹⁷

Data Analyses

Qualitative codes were transformed into dichotomous variables for quantitative analyses, frequency distribution, and cross-tabulation, to explore the relationship between each incident type and the respective contributory factors.24 All severe harm and death reports underwent a thematic analysis by Miss Rees and Dr Carson-Stevens to provide more in-depth insights into this subset of reports.20 Priority areas for improvement were identified based on frequently harmful incidents that also resulted in severe harm or death. Recommendations for improving these priority areas were informed by the factors contributing to them, systematic literature searches, and consultation with subject matter experts. Recommendations were articulated in an improvement tool called a driver diagram.25,26

Ethical Approval

Aneurin Bevan University Health Board research risk review committee waived the need for ethics review given the anonymized nature of the data (ABHB R and D reference number SA/410/13).

RESULTS

Baseline Characteristics

We included 1788 incident reports in the analysis after excluding 350 reports because they were about adult patients (n = 216), contained insufficient free-text information (n = 82), were reports about system issues that did not result in a patient safety incident (n = 32), or described deaths not associated with health care (n = 20).

Of the 1788 reports included, 763 described incidents in which children experienced harm, including 675 reports of low harm, 71 reports of moderate harm, 9 reports of severe harm, and 8 reports of death (see Table 1 for definitions). Incidents that resulted in severe harm and death will be referred henceforth as the "most harmful" incidents. Twelve parent incident types were evident, including those related to administration, medication, referral, and vaccination that were responsible for 71.8% of reports (see Fig 1 and Table 2). The most frequent contributory factors included documentation-related errors such as inaccurate medical records, resource issues such as staffing, staff cognitive issues (ie, mistakes), and clinical skills errors such as inadequate patient assessment (see Table 3).

Vaccine-related incidents were the most frequently described and contained the most descriptions of harm to patients (n = 472, 70.6%), including 459 cases of low harm and 13 cases of moderate harm. Administration of vaccines at the wrong time (n = 104), the wrong number of doses (n = 237), and the wrong vaccine (n = 154) were the most frequently described vaccination incidents. The key factors contributing to these incidents

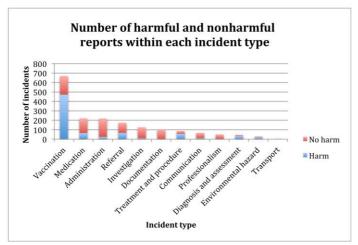


FIGURE 1
Twelve parent incident types and the proportion of harmful incidents described within each incident type.

included documentation errors (n = 129) and staff mistakes (n =101). However, vaccination incidents did not result in any cases of severe harm or death. We present a summary of findings from the most frequent and most harmful incident types that were described as resulting in severe harm or death. The 4 key incident types presented (diagnosis and assessment, treatment and procedures, referral issues, and medication provision) form the basis of 3 crosscutting priority areas requiring improvement (see Table 2 and Fig 2). Although administration incidents did result in severe harm and death, they were infrequently

harmful and were therefore not prioritized for inclusion here.

Diagnosis and Assessment-Related Incidents

Diagnosis and assessment incidents were described in 45 reports, and most (n = 34, 76%) described harm, including 4 deaths, 3 cases of severe harm, 11 cases of moderate harm, and 16 cases of low harm. The most harmful incidents frequently described delayed diagnosis or insufficient assessment of children with signs of child maltreatment, respiratory distress, diabetes, peritonsillar abscess, cervical fracture, and gastroenteritis during

TABLE 2 Twelve Parent Incident Types Presented With Their Frequency and the Proportion of Harmful Reports

Incident Type	Incidents						
	All, n	Harmful, n (%)	Severe Harm and Death, n				
Diagnosis and assessment	45	34 (76)	7				
Environmental hazarda	28	20 (71)	0				
Vaccination	669	472 (70.6)	0				
Treatment and procedure	86	57 (66)	2				
Transport ^a	4	2 (50)	0				
Referral	174	66 (38)	3				
Medication	221	65 (30)	2				
Communication	67	10 (15)	0				
Administration	219	24 (11)	3				
Investigation	126	10 (8)	0				
Documentation	99	2 (2)	0				
Professionalism	50	1 (2)	0				
Total	1788	763 (42.7)	17				

^a Rare (frequency <30)

an *Escherichia coli* 0157 outbreak (see examples 4.1 and 4.2 in Table 4). Potential contributory factors were described for 40% (n = 18) of incidents (Table 5). Knowledgerelated errors (n = 7), such as missing signs of child maltreatment (example 4.1) and errors of patient assessment (example 4.2) (triaging, history taking, examination, and investigation) (n = 7), were most frequently described.

Treatment and Procedure–Related Incidents

Most (n = 57, 66%) treatment and procedure incidents described harm to children; these included 1 death, 1 case of severe harm, 6 cases of moderate harm, and 49 cases of low harm. The most harmful incidents were often the result of incorrect treatment decisions, particularly treatment of asthma and diabetes (see examples 4.3 and 4.4 in Table 4). Potential contributory factors were described for 71% (n = 61) of incidents, including mistakes (n = 23)such as inadvertently getting glue in a child's eye when gluing a forehead laceration; poor-quality equipment (n = 13) such as a broken refrigerator: and poor knowledge (n = 10) such as how to correctly manage first presentation of diabetes in children (example 4.3, Table 5).

Referral-Related Incidents

Harm from referral-related incidents was described in 66 reports, including 2 deaths, 1 case of severe harm, 23 cases of moderate harm, and 40 cases of low harm (see examples 4.5 and 4.6 in Table 4). Referral decisions were frequently reported as harmful (n = 20, 59% of reports) and were often (n = 10,50%) preceded by incidents of diagnosis and assessment (example 4.5). Example 4.5 illustrates a failure to refer a child as a result of insufficient assessment and diagnosis, whereas example 4.6 illustrates a referral issue, which is not described as the result of insufficient

TABLE 3 Frequency of Reported Contributory Factors

Contributory Factors	Reports, n (%)
Documentation errors (inadequate handling or updating of patient records)	207 (11.6) ^a
Inaccurate patient records	135 (7.6)
Transfer of records	80 (4.5)
Staff mistakes (cognitive lapses including inattention and distraction)	149 (8.3)
Resource issues (inadequate supply of treatment, equipment, or staff)	145 (8.1) ^a
Human resources (staffing)	74 (4.1)
Equipment availability and safety	40 (2.2)
Treatment availability	31 (1.7)
Other resources	2 (0.1)
Failure to follow protocol (not adhering to organizational guidelines)	109 (6.1) ^a
Treatment protocol	44 (2.5)
Other and nonspecific protocols	37 (2.1)
Child protection protocol	21 (1.2)
Acutely unwell patients	7 (0.4)
Clinical skills errors (inadequate execution of routine tasks)	93 (5.2) ^a
Communication	70 (3.9)
Error in patient assessment	20 (1.1)
Other clinical skills	5 (0.3)
Treatment provision errors (issues with the process of medication delivery)	81 (4.5)
Medication names confused	43 (2.4)
Other	34 (1.9)
Patient names confused	4 (0.2)
Knowledge errors (insufficient knowledge or inadequate application of it)	53 (3.0)
Inadequate guidelines or protocols (existing guidelines not fit for purpose)	31 (1.7)
Poor continuity of care (issues with the coordination of services)	29 (1.6)
Total	897 (50.2)

^a Some reports contained descriptions of multiple types of a contributory factor.

diagnosis and assessment. Among the most harmful incidents, themes included failure to refer children with child protection concerns; decisions not to refer acutely unwell children with respiratory difficulties; and failure to follow up or refer children with development delays. In addition, a sudden unexplained death was reported in a child with a history of seizures for which there was no evidence of referral.

Potential contributory factors were described for 68% (n = 118) of referral-related incidents including errors of patient assessment (n = 13), failure to follow child protection protocols (n = 16, example 4.5), staffing issues (n = 26), documentation errors (n = 25), and poor knowledge (n = 18) (Table 5).

Medication-Related Incidents

Most (n = 156, 70.6%) medication-related incident reports did not describe harmful outcomes, although 2 reports described severe harm, 12 described moderate harm, and 51 described low harm. Of the incidents resulting in moderate or severe harm, prescribing or dispensing a medication overdose was a recurring theme. Two incidents involved furosemide overdose, and another involved

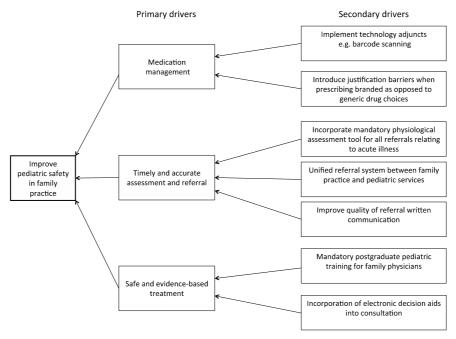


FIGURE 2Driver diagram demonstrating priority areas for improved pediatric care in family practice

Diagnosis and Assessment-related Incidents

Example 4.1: Mother of a 3-year-old noticed blood on child's pajamas. Physician diagnosed cystitis and prescribed antibiotics without examining child or taking a urine sample. Bleeding in a 3-y-old is unusual, and physician should have sought a pediatric review. Patient seen by pediatrician 5 d later when it was discovered that child had been sexually abused. (Severe harm)

Example 4.2: Child was wrestling and fell. Child screamed and neck appeared to lock. Out-of-hours physician examined child seated on mother's lap. Child was limited in his movements. Mother felt examination was not thorough. Physician advised the injury was not serious. Child was unable to walk to get to the toilet. Mother was concerned and rang out of hours (unscheduled care) service again. New physician said child was in shock and that he did not have a fracture because she would be able to feel it. On Sunday the child was still immobile and laying flat and complaining of pain. On Tuesday, mother contacted the family physician who believed the child had a serious injury. He has a spinal fracture between C2 and C3. The neck is currently at a 30-degree forward angle. (Severe harm)

Example 4.3: Patient saw family doctor with 3-week history of symptoms of diabetes. After documenting high blood sugar, she was started on an oral hypoglycemic drug. Brought to emergency department with blurred vision. Patient was inappropriately treated and not referred promptly. (Moderate harm) Example 4.4: Admitted to ICU with diabetic ketoacidosis. The incident is the result of advice given to mother by the family physician to omit evening insulin. (Severe harm)

Referral-Related Incidents

Example 4.5: Child seen for child protection medical on February 15, with facial bruising, described as being suggestive of possible slap mark and finger print bruising. On February 13, child was seen by 2 practice nurses for another matter who advised mother to take child to the out-of-hours family physician if bruising got worse or if the child became unwell. The serious significance of the bruising nor the need for a referral to be made did not seem to be considered. (Severe harm)

Example 4.6: Boy seen by family physician July 14 with history of increased thirst, polyuria, waking up in the night for micturition, lethargy, tiredness for 2 to 3 wks. Blood tests (July 17) showed high blood sugar (13.9 mmol/L). Notes (July 19) say to review and consider if he needs to be referred. July 20, child was seen with sugar and ketones in urine and referred to hospital in ketosis. Any child with suspected diabetes should be referred within 24 h to hospital pediatrician. (Moderate harm)

Medication—related Incidents

Example 4.7: Risperidone dose (5 mg twice daily) recommended by child and adolescent mental health team for a child that they had assessed. The family doctor requested a consultant's opinion and was informed that this was the dose agreed by the team. Patient later presented with signs of toxicity (dyskinetic tongue). (Severe harm)

Example 4.8: Dispensed [acetaminophen] elixir instead of lactulose—similar products stored together in same-colored containers. (Low harm)

a 10-fold overdose that resulted in prolonged hospitalization of the child. Another report described a risperidone overdose prescribed by a family physician on the instruction of a child psychiatrist (example 4.7).

Table 4 illustrates examples of medication-related reports. Most medication incidents (n = 119,

TABLE 5 Frequency of Contributory Factors Described for Each Key Incident Type

Contributory Factors	Incident Type, n								
	Diagnosis and Assessment	Treatment and Procedure	Referral	Medication					
Documentation errors	1	2	40	4					
Staff mistakes	0	23	1	17					
Resource issues	3	21	28	7					
Failure to follow protocol	4	4	28	6					
Clinical skills errors	9	6	26	15					
Treatment provision errors	1	0	0	24					
Knowledge errors	7	10	16	8					
Inadequate guidelines or protocols	0	4	11	1					
Poor continuity of care	1	1	6	3					
Total	26	71	156	85					

Note: some incidents had multiple contributory factors.

53.8%) were prescribing errors (see Table 6). Potential contributory factors were described for 33% of those reports (n = 73). Errors in the process of treatment provision (n = 24), such as confusing medication names (n = 12), were frequently described. Communication errors (n = 10, example 4.7) and staff mistakes, such as misreading medication names (n = 17, example 4.8), were also frequently described (Table 5).

Driver Diagram

Three broad crosscutting priority areas (denoted "primary drivers") with more specific articulations of the priority concepts (denoted "secondary drivers") were identified as candidate areas eligible for development and testing of interventions in practice to improve pediatric care (see Fig 2). The driver diagram was authored collaboratively by all study authors after analysis.

DISCUSSION

Of the harmful incidents reported from family practice, key pediatric safety issues include timely diagnosis, assessment, and referral of unwell children to the hospital; reliability of safe, evidence-based treatment; and prescribing and dispensing of medications. These priority areas include the 4 most harmful and frequently reported incident types. We recommend that improvements in these areas focus on the factors frequently reported to contribute to them. Examples are included in Fig 2 as a driver diagram that family practice teams could use as the basis for planning their own improvement efforts, supplemented by insights from their own locally available patient safety data 15

This is the first analysis of a large volume of pediatric incidents in family practice from a national database. Rigor was improved by careful consideration of methods

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TABLE 6 Subtypes of Medication Incidents Reported

Medication Incident Subtypes	Reports,			
Prescribing	119			
Dispensing	53			
Administering	25			
Adverse reaction	9			
Other	15			
Total	221			

before analysis, double-coding, weekly meetings to discuss coding, and keeping an audit trail to aid reflexivity.^{20,27}

Reporting systems rely on data input (reporting) to generate learning; unfortunately, only 50% of reports included in this analysis described potential contributory factors, and reporters do not routinely describe the root-cause, system-level factors contributing to incidents. In addition to variable data quality, underreporting is a wellacknowledged issue with the NRLS.28 This affects the validity and generalizability of our findings, which should be regarded as essentially inductive, and hypothesis generating, requiring confirmation in further studies.^{29,30} Despite this, reporting to the NRLS has increased in the last decade, providing large quantities of data from which to generate learning.31,32 There may be other harmful incident types occurring in family practice that are underreported owing to fear of reprimand.33 However, despite limitations from underreporting and reporting biases, analyses of NRLS data have played an important role in generating lessons to mitigate harmful incidents in other areas of clinical practice.34,35 Incident report data offer a single lens on patient safety, and our findings must be interpreted cautiously alongside other data sources.

The relatively low number of reports describing diagnosis and assessment errors was unexpected, as family practice has been widely criticized for failures in recognizing sick children.^{11,36} For example,
Thompson et al found that 50% of
children with meningococcal
infection are sent home from their
first consultation.³⁷ Such failures
likely contribute to the United
Kingdom's relatively high child death
rates from meningococcal disease,
pneumonia, and asthma, where
first-access services such as primary
care are integral.^{38,39} Diagnosis and
assessment incidents tend to be
a reflection of providers' skills and
competencies and may be less likely
to be reported.

Management of long-term pediatric conditions, such as epilepsy, asthma, and diabetes, has been repeatedly cited as requiring improvement. 9,11,36,39 For example, most (75%) pediatric asthma-related hospital admissions are thought to be preventable with better primary care. 40 Our findings reflect the significant potential for harm to children that exists as a result of poor management and treatment of long-term pediatric conditions in primary care.

Referrals made to hospitals, management of long-term conditions, and diagnosis and assessment of acutely ill children were all identified as problem areas, for which training interventions could be a generic improvement strategy. The numerous reports describing referral issues were anticipated, as UK primary care acts as a gateway to specialist hospital services. However, ~36% of referrals to pediatricians are thought to be avoidable with better family physician training.39,41 Currently, as few as 40% of family physicians have pediatric training at a postgraduate level, causing many to support mandatory family physician pediatric training.39

Our findings support such calls for mandatory pediatric training, as reported harm from incidents related to referral, treatment, diagnosis and assessment were frequently associated with errors of knowledge and patient assessment.39 Training should be accompanied by regular measurement and monitoring of physician performance. Improvement in the timely referral of unwell children should focus on simplifying referral protocols for children at high risk, making them easier to follow and giving them face validity; for example, using an electronically generated and transmitted unified referral form. This could be supplemented with online decision support for physicians' decisions regarding the management, diagnosis, assessment, and referral of unwell children.42

High numbers of reports describing harmful medication incidents were expected, as they are often harmful and widely cited as the most common medical errors.^{1,43–46} In addition, medication incidents in pediatrics may be ≤3 times more common than in adults.43,47 Most medications are prescribed off license and can require complex dose calculations.48,49 Pediatric medication incidents (including their extent and methods to reduce them) have been extensively explored in the hospital setting, but less so in family practice.43,47,50-54 Our findings emphasize the importance of verification procedures and support barcode scanning of medications during dispensing and using generic medication names (rather than brand names) to reduce mistakes from inattention or distraction and communication errors.55-57

CONCLUSIONS

This is the first analysis of nationally collected pediatric safety incidents from family practice. Potential priority areas for improved care for children in this setting have been identified in timely diagnosis, assessment, and referral of unwell children; reliable delivery of safe treatment; and medication provision. Suggested

improvements in these areas include mandatory pediatrics training for all family physicians; use of electronic tools to support diagnosis, management and referral decision-making; and, use of technological adjuncts such as barcode scanning systems for medication provision.

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POTENTIAL CONFLICT OF INTEREST: Dr Carson-Stevens and Professor Edwards are co-chief investigators, and Drs Panesar, Professor Sheikh, Dr Mayor, Professor Avery, and Professor Donaldson are co-applicants, of a NIHR HS&DR grant to characterize patient safety incident reports in primary care. Professor Donaldson was the Chairman of the National Patient Safety Agency (NPSA) (2010 to 2012) and is currently involved in the program of research associated with the National Reporting and Learning System (NRLS); he is also the World Health Organization's Patient Safety Envoy. Dr Panesar is a former clinical adviser at the NPSA (2008 to 2010), then special adviser to Professor Donaldson (2010 to 2012), and a former academic clinical fellow at Imperial College London working for the NRLS research program.

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1.5 Lancet commentary (iatrogenic harm in children)

Carson-Stevens A, Edwards A, Panesar S, Parry G, Rees P, Sheikh A, Donaldson L. Reducing the burden of iatrogenic harm in children. Lancet. 2015 Apr 25;385(9978):1593-4. doi: 10.1016/S0140-6736(14)61739-6.

Reducing the burden of iatrogenic harm in children

In 2013, as many as 6.3 million children worldwide died before their fifth birthday.1 Children have an increased risk of health-care-related harm because of factors including the complexity of prescribing and dispensing of drugs, a reduced physiological reserve compared with adults, and dependency on others (ie, parents and health-care providers) to recognise the emergence of a hazardous situation.² Despite these factors, little research has been done of the contributions of substandard care and iatrogenic harm to deaths in childhood. Furthermore, health-care-related harm has featured little in the Millennium Development Goals; the Sustainable Development Goals must include this issue, and establish priority to resolve the existing paucity of research related to children and young people and promote policy making for strengthening of healthcare systems to cater for children's needs.3,4

Confidential enquiries into child deaths in the UK showed that 26% of deaths had identifiable failures in care, and a further 43% had potentially avoidable factors. ⁵⁶ In the USA, an estimated 15–35% of children admitted to hospital have health-care-associated harm. ⁷ Despite patient safety being an established health policy concern for adults, little has been done to estimate the frequency of harm and its associated burden in the care of children.

Research in adults has shown that a range of methods can be successfully used to study substandard care and iatrogenic harm. Trigger tools can identify, within medical records, features suggestive of adverse events. Safety indices can be constructed from administrative data to estimate what incidents happen, whilst methods such as root cause analyses and incident reporting systems can aid understanding of why incidents happen.²⁸ This work now needs to be extended to children and young people. That said, collection, analysis, and use of such data in health care remain problematic, with few health systems showing that they can reduce risk for future patients.

In the England and Wales National Reporting and Learning System, more than 65 000 reports of safety incidents involving children are received every year; about 30% of those reports record incidents that resulted in harm or death to children (Carson-Stevens A, unpublished). Useful lessons can be learned from these reports: 77 safety notices were issued between 2004 and 2012 by the National Patient Safety Agency

covering the UK National Health Service in England and Wales, including recommendations for keeping babies with a family history of medium-chain acyl-CoA dehydrogenase deficiency safe in the neonatal period, for prevention of harm to children from parents with mental health problems, and for nasogastric placement in infants. Themes generated from incident reports can help in the design of logic models that can inform quality improvement interventions. Moreover, problems that would be overlooked at a local level because of small numbers can become evident when aggregated across wider areas.

Globally, a systematic approach is needed to generate data of this nature and to construct basic metrics of harm in children and young people, and to ensure that these data are reflected within international development goals. The Global Burden of Disease study¹ could provide annual estimates of the scale of iatrogenic harm and could inform health policy makers about where to direct efforts to mitigate such harm. WHO has proposed a Minimal Information Model to provide a dataset in all countries for sharing of patient safety incident reports.¹¹O This model could improve understanding of the nature, and particularly the causes, of unsafe care. We believe that present incident reporting systems are undervalued and underused, garnering little respect from the health information and research communities.

An international patient safety learning system is needed, designed to describe care failures or safety incidents, shape priorities for improvement, corroborate



insights from research studies, develop potential solutions for application in practice, and share learning of the context-specific approaches of application of solutions. In view of the different maturity levels of data collection across different economic settings, the system would benefit from a phased effort to develop the necessary infrastructures for data sharing in each country. Member countries of such a system must hold autonomy to undertake their own searches informed by insights gleaned from surveillance of their own national safety data. Moreover, countries need to develop a culture of open reporting from staff, parents, and patients to provide future high-quality incident reports, build improvement capability within their workforce to apply the lessons learned, and educate family and patients for early warning and early action to mitigate care failure. Such development will require a combined enterprise between clinical and patient safety experts and the commitment of health leaders and policy makers.

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Hip fractures: comprehensive geriatric care and recovery

Published Online February 5, 2015 http://dx.doi.org/10.1016/ 50140-6736(14)61592-0 See Articles page 1623 Hip fractures are a worldwide public health issue, with devastating consequences for both patients and their families. In 2000, 1·6 million hip fractures were reported worldwide¹ and epidemiological studies estimate a 12·1% and 4·6% lifetime risk of hip fracture for women and men, respectively.² Clinical and social consequences of hip fracture include death, depression, fear of falling, disability, institutionalisation, and social isolation.³ For old people, the ability to remain mobile is an essential aspect of quality of life, and is crucial for the preservation of independence. Many older patients develop disabilities in mobility after hip fracture surgery, and more than 30% do not regain independent ambulation

1 year later.⁴ A better understanding of this suboptimum recovery, and innovative, effective interventions beyond surgery, are clinical and public health priorities.

In *The Lancet*, Anders Prestmo and colleagues⁵ present their well designed study providing strong evidence that, for patients aged 70 years or older with hip fractures, perioperative comprehensive geriatric care given in a dedicated ward improves short-term and long-term function in mobility, as compared with treatment in a traditional orthopaedic care unit. 198 patients were randomly assigned to comprehensive geriatric care, and 199 to usual orthopaedic care. Mobility was assessed by the trial's primary endpoint,

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1.6 Vaccine manuscript (vaccination errors in primary care)

Rees P, Edwards A, Powell C, Evans HP, Carter B, Hibbert P, Makeham M, Sheikh A, Donaldson L, Carson-Stevens A. Pediatric immunization-related safety incidents in primary care: A mixed methods analysis of a national database. Vaccine. 2015 Jun 26. pii: S0264-410X(15)00869-5. doi: 10.1016/j.vaccine.2015.06.068.

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Pediatric immunization-related safety incidents in primary care: A mixed methods analysis of a national database

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ABSTRACT

Background: Children are scheduled to receive 18–20 immunizations before their 18th birthday in England and Wales; this approximates to 13 million vaccines administered per annum. Each immunization represents a potential opportunity for immunization-related error and effective immunization is imperative to maintain the public health benefit from immunization. Using data from a national reporting system, this study aimed to characterize pediatric immunization-related safety incident reports from primary care in England and Wales between 2002 and 2013.

Methods: A cross-sectional mixed methods study was undertaken. This comprised reading the free-text of incident reports and applying codes to describe incident type, potential contributory factors, harm severity, and incident outcomes. A subsequent thematic analysis was undertaken to interpret the most commonly occurring codes, such as those describing the incident, events leading up to it and reported contributory factors, within the contexts they were described.

Results: We identified 1745 reports and most (n = 1077, 61.7%) described harm outcomes including three deaths, 67 reports of moderate harm and 1007 reports of low harm. Failure of timely vaccination was the potential cause of three child deaths from meningitis and pneumonia, and described in a further 113 reports. Vaccine administration incidents included the wrong number of doses (n = 476, 27.3%), wrong timing (n = 294, 16.8%), and wrong vaccine (n = 249, 14.3%). Documentation failures were frequently implicated. Socially and medically vulnerable children were commonly described.

Conclusion: This is the largest examination of reported contributory factors for immunization-related patient safety incidents in children. Our findings suggest investments in IT infrastructure to support data linkage and identification of risk predictors, development of consultation models that promote the role of parents in mitigating safety incidents, and improvement efforts to adapt and adopt best practices from elsewhere, are needed to mitigate future immunization-related patient safety incidents. These priorities are particularly pressing for vulnerable patient groups.

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Abbreviations: NRLS, National Reporting and Learning System; WHO, World Health Organization.

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1. Introduction

Each immunization represents a potential opportunity for immunization-related safety incidents, and given their prevalence and potential preventability the World Health Organization (WHO) has recognized this as a priority area for healthcare improvement [1]. Without address, population-level protection could be compromised, national campaign efforts to promote uptake hindered, and the trust of patients and families reduced [1,2].

Reviews of patient medical records estimate 27–35% of immunizations involve an error [3–6]. Analysis of patient safety incident reports, such those within the US MedMARx reporting system, have suggested additional doses, the wrong immunization, and the wrong dose given are the main types of errors involved [7]. Further, surveillance systems collecting information about adverse drug reactions, such as the Vaccine Adverse Event Reporting System, have also identified common sources of errors, but they primarily focus and receive reports on the health problems, illnesses or symptoms experienced by patients following immunization [8,9]. Despite those insights, analyses of such reports are often limited by the lack of comprehensiveness of reports.

Whilst previous studies have classified immunization-related safety incidents in terms of their type [7,9,10], few studies have comprehensively explored the underlying factors contributing to them [3–5,7–11]. Incident reporting systems, like the England and Wales National Reporting and Learning System (NRLS)², include a range of patient safety incident types, specifically descriptions of, "any untoward incident that may or may not have led to harm" and invite free text descriptions of what happened, perceived potential contributory factors and actions to prevent future occurrence. Examination of report content can support the design of learning interventions to mitigate future events [12,13].

We aimed to characterize the nature and severity of immunization-related patient safety incidents involving children in primary care in England and Wales, in order to: identify priority issues for improvement; and generate hypotheses about change ideas that could form the basis of improvement recommendations and interventions.

2. Method

2.1. National reporting and learning system

The NRLS² is a national reporting system that collates locally generated reports that are received from healthcare organizations throughout England and Wales. The NRLS was launched in 2003 and receives approximately 100,000 reports a month, the majority written by healthcare professionals. It receives over 65,000 reports per annum involving children [13]. Reporting patient safety incidents that resulted in severe harm or death of a patient became mandatory in June 2010; however, before this all reporting was voluntary, and reporting remains voluntary for incidents resulting in no, low, or moderate harm [14,15].

Each report contains categorical information about location, patient age, incident type, and reporter perception of harm severity – collected in a structured report form – as well as free-text descriptions of the incident [14,15].

2.2. Sample selection

Over 20,000 pediatric, primary care-related incidents are present in the NRLS. Immunization-related safety incidents were identified by free-text key term searches, which we have found to generate both a sensitive and pragmatic means for identifying reports. Search terms and their permutations were

informed by brand and generic vaccine names from the British National Formulary and by an analysis of a pilot sample of over 600 immunization-related reports [10,16]. We included any immunization-related safety incidents occurring in primary care involving a child aged under-18 years between 2002 and 2013.

2.3. Data coding

Incident report free text was read and codes were applied to describe incident type, potential contributory factors, harm severity, and incident outcomes. The 'Recursive Model of Incident Analysis' method was used as a set of rules for applying codes in a chronological order (Supplement 1) [17]. This permitted modeling of the sequence of events leading to the primary incident type that resulted in the outcome experienced by the patient (Supplement 2). A 'severity of harm' had already been assigned to each report by the reporter; based on our interpretation of the report content, the severity of harm was re-classified using definitions from the WHO International Classification of Patient Safety when required (Table 1) [18]. Generic and brand names of vaccines were recorded. Incident types were independently double coded for a random 20% sample of reports (PR and HPE) with 3rd person arbitration (ACS) when necessary.

2.4. Data analysis

Relationships between codes were examined using frequency distributions and cross-tabulations. Two-way cross-tabulations were generated between age, incident type, vaccine type and contributory factors, compared to harm. Associations between these were examined using the Fisher's exact test. To adjust for confounding of vaccine type on the effect of incident type on harm severity, a Mantel Haenszel adjusted approach was taken by stratifying vaccine and incident type.

2.5. Thematic analysis

Reports that contained the most frequently occurring incident type or contributory factor codes were re-read to generate additional insights and interpretations about those incidents and the kind of contexts in which they occurred [12]. Qualitative data software (NVIVO 9, QSR International) was used to independently analyze the reports and create new codes to capture this information. Themes that helped support our understanding of those incidents, and why they might have occurred, were agreed between PR and ACS [12,19].

2.6. Ethical approval

Aneurin Bevan University Health Board research risk review committee waived the need for ethical review given the anonymized nature of these data (ABHB R and D Ref number: SA/410/13).

3. Results

3.1. Characteristics of reports

Free-text searches identified 2298 reports, of which 1745 were included. Reports were excluded if they: were not immunization-error related (n=464), e.g. describing a child who had received appropriate immunizations; contained insufficient free text (n=24); or described issues that did not result in a patient safety incident (n=65). Reports were submitted from 254 NHS providers in England and Wales, including 1052 reports from community nursing and 596 reports from general practice settings. Cohen's

Table 1The vaccines involved in reported incidents and the number of reports they are involved in (note: *2 vaccines were involved in 1 child's death).

Vaccine	No harm	Low harm	Moderate harm	Death	Frequency (%)
Measles Mumps Rubella (MMR)	86	269	6	_	361 (18.2)
Pneumococcal conjugate (PCV)	140	160	4	3	307(15.5)
Diphtheria tetanus acellular pertussis/inactivated polio/haemophilus influenza type B (DTaP/IPV/Hib)	97	142	1	1	241 (12.1)
Meningitis C (Men C)	103	122	1	_	226(11.4)
Haemophilus influenza b/meningitis C (Hib/Men C)	70	123	2	-	195(10)
Diphtheria tetanus acellular pertussis/inactivated polio (DTaP/IPV)	29	145	1	-	175(9)
Human papilloma virus (HPV)	52	59	14	-	125 (6)
Bacillus Calmette-Guérin (BCG)	42	52	7	_	101(5)
Tetanus, diphtheria and inactivated polio (Td/IPV)	15	65	13	-	93(5)
Haemophilus influenza b (Hib)	13	37	-	-	50(3)
Hepatitis B	24	10	7	_	41(2)
Hepatitis A	10	11	0	-	21(1)
Influenza	11	8	1	_	20(1)
Other	6	9	1	-	16(1)
Rotavirus	5	2	_	-	7(0)
Typhoid	2	4	-	_	6(0)
Total	705	1218	58	4*	1985(100)

Definitions of Harm: No harm—patient outcome is not symptomatic and no treatment is require; Low harm—patient outcome is symptomatic, symptoms are mild, loss of function or harm is minimal and intermediate but short term, and no or minimal intervention is required; Moderate harm—patient outcome is symptomatic requiring intervention, an increased length of stay, or causing permanent or long-term harm or loss of function; Severe harm—patient outcome is symptomatic, requiring life-saving intervention or major surgical/medical intervention, shortening life expectancy or causing major permanent or long term harm or loss of function; and, Death—on balance of probabilities, death was caused or brought forward in the short term by the incident.

kappa statistic of inter-rater (coding) reliability was high, k = 0.77, n < 0.001

Most reports involved children aged less than three years old (n=952, 55%) and peaks in frequency occurred, as expected, within the age groups children receive most vaccinations. Tables 1 and 2 highlight the most commonly cited vaccines and their associated harm severity outcomes. Most reports (n=1135, 65%) described outcomes, which included: patient inconvenience (n=801, 45.9%) such as receiving unnecessary treatment (n=481, 27.6%) and requiring additional treatment (n=379, 21.7%); clinical patient harm (n=205, 11.7%) such as injuries (n=72, 4%); and exposing the patient to risk (n=139, 8%), for example by leaving them vulnerable to immunization preventable diseases (n=108, 6%).

Administration was the most frequent incident type (n = 1282, 73.5%), with 'wrong number of doses', 'wrong timing', and 'wrong vaccine' described in (n = 1019, 79.5%) of those reports (Table 2). These are analyzed in further detail below.

There was a strong association between the type of administration-related incident and harm (p < 0.001) and the type of vaccine and harm (p < 0.001). After stratification into both 'incident' and 'vaccine' strata there was an association between type of administration incident and harm (in all vaccine strata p < 0.001), thus vaccine type did not explain the varied risk of reported harm.

3.2. Administration of the wrong number of doses

Children who received the wrong number of doses were typically under six years old, most frequently (less than two years old n=141,30%), and typical vaccines implicated included Measles Mumps Rubella (MMR) (n=156,33%), Diphtheria Tetanus acellular Pertussis/Inactivated Polio (DTaP/IPV) (n=87,18%), and Haemophilus influenza b/Meningitis C (Hib/Men C) (n=77,16%) (Table 4). Reports frequently described harm (n=448,94.1%), typically because the child received unnecessary additional

vaccinations—a low harm event (Example 3.1, Table 3). One incident resulted in an adverse reaction necessitating a hospital admission—a moderate harm. There was no evidence to suggest association between vaccine type and harm (p=0.49) within this stratum (wrong number of doses).

Such incidents were frequently the result of prior incidents involving documentation failures (n = 188, 40%), including documentation not being up to date (n = 128, 27%), available (n = 45, 9%), or accurate (n = 15, 3%) (Examples 3.1–3.3, Table 3; Supplement 2). Additional associated incidents included administration of the wrong vaccine (n = 49, 10%), communication failures (n = 26, 5%), and difficulties making appointments (n = 19, 4%).

Table 5 presents the contributory factors described. Patient and parent contributory factors included: inadequate knowledge (n = 40, 8%), for example not being aware of which vaccines were needed or had previously been received; being new to the area or family practice (n = 36, 8%); and eight incidents (2%) were partly the result of a child being in 'out-of-home' care (e.g. foster care) (Examples 3.2 and 3.3, Table 3). Staff factors included: not following a protocol (n = 60, 13%), for example not checking the medical records before administration; and mistakes (n = 67, 14%) such as misreading vaccine names (n = 10, 2%).

3.3. Administration at the wrong time

Vaccines administered at the wrong time (n = 294, 16.8%) were typically described as deviating from the recommended national immunization schedule. Most children were aged under 1 year old (n = 175, 60%). Vaccines typically involved were Pneumococcal conjugate (PCV) (n = 91, 31%), Diphtheria Tetanus acellular Pertussis/Inactivated Polio/Haemophilus influenza type B (DTaP/IPV/Hib) (n = 83, 28%), and the MMR (n = 48, 16%) (Table 4). There was no evidence to suggest an association between vaccine type and harm (p = 0.51) within this stratum (wrong time).

Of 55 (19%) harmful incidents, eight incidents resulted in moderate harm—typically these reports described delayed prophylactic

 Table 2

 The frequency and severity of harm described for each age group and for each primary incident type (N.B. this table does not include the frequencies of contributory incidents).

	Severity of harm								
	No harm	Low harm	Moderate harm	Death	Frequency (%)				
Age									
Under 28 days	15	7	3	-	25(1)				
1 month to 1 year	307	246	13	3	569 (32.6)				
2 to 4 years	206	433	9	_	648 (37.1)				
5 to 11 years	41	133	5	-	179 (10)				
12 to 17 years	99	188	37	-	324 (18.6)				
Incident type					8 9				
Vaccination									
Administering	438	824	17	3	1282 (73.5)				
Wrong number of doses	28	447	1	=	476				
Wrong timing	239	44	8	3	294				
Wrong vaccine	97	150	2	_	249				
Not administered	14	65	1	12	80				
Wrong dose	26	31	-	-	57				
Expired vaccine	3	46	_	_	49				
Non-specific	10	13	=	200	23				
Contraindicated vaccine	14	5	1		20				
Wrong patient	2	14		_	16				
Wrong site	1	2	3	_	6				
Used needle	3	2	-	-	5				
Wrong storage	_	4	_		4				
Wrong route	1	1	1	-	3				
Adverse reaction	-	103	43	-	146 (8)				
Prescribing and dispensing	26	_	_		26(1)				
Reconstitution error	-	4	_		4(0)				
Batch recall	4	=	_	_	4(0)				
Insufficient supply	i	223	24		1(0)				
Non-specific	î	_	_	(-	1(0)				
Documentation	97	5	1	-	103 (6)				
Records not up to date	62	3	1	12	66				
Records inaccurate/unclear	29	2	_		31				
Record availability	6	-	-		6				
Administration	50	4	2	_	56(3)				
Appointment management	36	3	-	-	39				
Transfer of information	12	1	2	_	15				
Other	2	_	_		2				
Procedural skills	4	53	_		57 (3)				
Communication	44	9	_	_	53 (3)				
With patients or parents	42	9	_		51				
Between HCPs	2	_		_	2				
Other	3	5	4	12	12(1)				
Total	668	1007	67	3	1745 (100)				

administration of Hepatitis B and Bacillus Calmette-Guérin (BCG) vaccines to high-risk newborns. Three other incidents of delayed vaccination-contrary to the national recommended schedule-were potentially implicated in child deaths from meningitis and pneumonia, (Example 3.4, Table 3). Reports describing low harm outcomes typically described children who required additional vaccinations for adequate immunity. Timing-related vaccine incidents were preceded by incidents involving: appointment management (n = 60, 20%), documentation failures (n = 31, 11%), communication failures (n = 21, 7%), transfer of documentation (n = 16, 5%), and other vaccine-related incidents such as administration of the wrong vaccine (n = 12, 4%) (Supplement 2).

Table 5 highlights contributory factors implicated in these incidents such as: failure to follow protocol (n=32, 11%); poor continuity of care (n=22, 7%), for example community nurses not receiving birth notifications from secondary care; and staff mistakes (n=22, 7%) (Examples 3.4–3.7, Table 3).

3.4. Administration of the wrong vaccine

Administration of the wrong vaccine was described in 249 reports (14.3%) and they were largely given to children under one year old (n=121, 49%). Of these, 152 (61%) incidents resulted in harm, typically because children received unnecessary vaccinations and required additional treatment i.e. the vaccine that was

originally required. For example reports described administration of a Hib/Men C combination vaccine (n=31, 12%) when a single (non-combination) Meningococcal C (Men C) vaccine was scheduled. Men C (n=87, 35%), PCV (n=85, 34%), and Hib/Men C (n=61, 25%) vaccines were frequently implicated (Table 4). These incidents were rarely preceded by other incidents; however 18 (7%) resulted from documentation failures, and six (2%) resulted from immunizing the wrong child (Example 3.8, Table 3; Supplement 2). Within this stratum (wrong vaccine) there was strong evidence for an association between vaccine type and harm (p<0.001).

Contributory factors described were: staff issues which included mistakes (n=87, 35%) such as confusing vaccines with similar names or appearances (n=37, 15%) and failure to follow protocols (n=35, 14%), such as concurrently preparing vaccines for multiple children (Example 3.9, Tables 3 and 5).

3.5. Overarching themes-Responsibility and vulnerability

Many reports implied parents were partly responsible for the occurrence of incidents, for example if the parent did not bring parent-held records to appointments or did not provide accurate medical histories (Example 3.3, Table 3). An underlying expectation that parents should be aware of their child's immunization needs and history was apparent. However parents appeared to have similar expectations of, and confidence that, healthcare professionals

Table 3 Edited extracts from incident reports.

Wrong number of doses

Example 3.1: Patient presented with **stepmother** for pre-school booster. Written **consent** from **father** was brought but **parental held record** was **not available**. Nurse explained she was giving repevax and **MMR**. The following day stepmother called expressing concern that **MMR** had **already** been **given** in 2004. **Incomplete documentation** of initial dose of MMR booster. (Low harm)

Example 3.2: Child placed with adoptive parents who were advised by Social Worker to attend physician to complete primary vaccinations. Attended surgery with parental held records but no family practice records were available. Only two immunizations were recorded in the parental held record. Immunization given with consent. Later informed by social services that child had already completed her primary immunizations. Family practice records checked and confirmed above. (Low harm) Example 3.3: Patient received the third primary immunizations twice in error once in Ghana and once at the health center. Mother failed to notify the health visitor of the first immunization. (Low harm)

Wrong timing

Example 3.4: An infant died from a streptococcal pneumonia—which could have been prevented if the child had received childhood immunizations. The mother stated she was not aware that her child should be immunized and the child was not registered at a family practice until *** Identified areas of concern include: the management of the child immunization processes, family practice registration processes and notifying child health of non-registered patients. (Death)

Example 3.5: Patients mother rang after receiving birthday card for child and realizing that her daughter had **not received** an **immunization appointment** since **transferring into the area**. Child health still had old address and old family practice details. Mother told to book immunization appointments with new family practice. (No harm)

Example 3.6: Patient's relative contacted health visitors regarding her child's immunizations, she reported she had **not received** any **appointments** for her child's third primary immunizations. **Child health** computer had **recorded wrongly** that the child had his third immunizations on the same day as he had his second immunizations. The patient **received** his third primary **immunizations late** because of this. Child health would not have been aware of this if the **parent** had not **contacted** the **service**. (No harm)

Example 3.7: Patient was scheduled for Hib/Men C vaccine, staff checked his immunization record and became aware that he already had this immunization. At this point I made an error. I told the mother that we could give the MMR/Prevenar. Mother's English is not perfect and she agreed. As I came to record the immunization, I realized my error there was a two-week gap between immunizations not 4. (No harm)

Wrong vaccine

Example 3.8: I was covering an immunizations session where normally there are 2 immunizers, the 2nd immunizer was sent home sick ten minutes prior to the start of the session. The healthcare assistant went to get the child and the mother for the immunization. The wrong immunization was administered to the child as we had the paperwork for a different child. The error was realized immediately, the child received Hib/Men C and MMR instead of 5 in 1 and Men C. (Low harm)

Example 3.9: Mother took her five-month-old baby to her family physician for his second DTaP/IPV/Hib vaccination. The staff nurse administered the wrong injection because she did not consult his medical records. The baby was given an MMR vaccination that should not be given until he is 13 months old. Staff Nurse says she was distracted during the appointment. The Nurse will re-train and demonstrate her competency through supervision. (Low harm)

Vulnerable child

Example 3.10: Baby received routine immunizations in baby clinic. The baby's consultant had written to the family physician requesting that the baby **not receive**immunizations until he is **one-year post chemotherapy. Mother believed** baby **should have immunizations** as he had received two flu vaccines between two sessions
of chemotherapy previously. Family physician had seen this letter and had put a note on baby record but the **note was not dated** and believed to be an automatic warning.
(No harm)

should be aware which immunizations their child required (Example 3.7, Table 3).

Collaboration between healthcare professionals and parents was reported as preventing some near-miss immunization-related safety incidents. There were reports detailing how parents had mitigated harm and prevented incidents by advocating for their children, chasing appointments, informing staff when the wrong vaccine was prepared, or reminding staff when a vaccine was contraindicated (Examples 3.5 and 3.6, Table 3). Lack of this parental safety net could be a factor in the immunization-related incidents described in children in 'out-of-home' care.

Socially vulnerable children such as those in 'out-of-home' care, and children of immigrant or traveller families, were described as experiencing incidents as a result of difficulty accessing care, poor continuity of care, and documentation failures. Accessing appropriate care was difficult for those vulnerable children for a range of reasons, including language barriers and inadequate parental knowledge of the need for vaccination or how to access primary care services, and in one case a child died (Example 3.4, Table 3).

Medically vulnerable children such as pregnant adolescents, those at risk of TB, Hepatitis B or HIV, or immunocompromised children, who required additional vaccinations or had contraindications to certain vaccines, were commonly represented in these reports. Numerous factors featured across those incident reports including: communication failures with children and parents; non-disclosure of medical conditions; non-adherence to the advice that parents were given by pediatricians; and poor staff and parent knowledge of vaccine contraindications in certain medical conditions (Example 3.10, Table 3).

4. Discussion

This study has shown that the types of immunization-related safety incidents experienced by children are consistent with studies undertaken in other countries. Safer immunization is a priority area for quality improvement, which should focus on administering the correct vaccine, the correct number of times, and at the correct time for all children, and the timely and correct immunization of

Table 4 the vaccines involved in the most frequently reported incident types and their frequencies.

	Vaccines														
Type of administration incident	MMR	PCV	DTaP/ IPV/Hib	Men C	Hib/Men C	DTaP/IPV	HPV	BCG	Td/IPV	Hib	Hepatitis B	Other	Hepatitis A	Influenza	Total
Wrong number of doses	156	75	63	49	77	87	8	20	36	16	_	5	3	1	596
Wrong timing	48	91	83	37	29	7	16	14	2	2	21	6	1	-	357
Wrong vaccine	41	85	56	87	61	49	8	3	23	1	1	5	6	4	430
Not administered	12	2	5	13	4	3	1	6	1	28	1		1	-	77
Wrong dose	12	1	1	7	3	1	2	9	-	-	5	1	5	3	50
Expired vaccine	12	_	1	2	4	7	1	5	1	3	1	1	2	2	42
Contraindicated vaccine	2	1	-	1	-	2	_	5	1	_	_	2	-	2	16
Wrong patient	7	1	4	2	1	1	2	-	1	-	-	-	-	-	19
Other	2	5	2	3	4	1	2	5	_	_	2	3	_	2	31

Table 5Frequency of described contributory factors (note: *some reports described more than one type of mistake).

Patient/parent factors Patient/parent behavior—the way in which patients or parents act or conduct themselves Non-compliance 60 Non-disclosure 12 Other 1 Patient/parent geography—the area where patients live New to area 62 Access difficulties 2 Patient health—factors relating to the patient's physical and mental wellbeing Allergy 22 Non-specific 4 Disability 4 Immunocompromised 3 Abnormal coagulation 2 Pregnancy 2 Patient/parent knowledge—insufficient 48 (5) knowledge Out-of-home care—children not in the care of their parents e.g. in foster care Patient/parent language—patient or parent unable to communicate in English Staff factors Mistake—cognitive lapses 240 (2000)	
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Distraction 22	
Misreading 18	
Inattention 10	
Similar patient names 9	
Failure to follow protocol—not adhering to 186 (2)	0)
organizational guidelines	
Knowledge—insufficient knowledge or 19 (2)	
inadequate application of knowledge	
Fatigue/stress—extreme tiredness, mental or 5 (1)	
emotional strain	
Other factors 3 (0)	
quipment/vaccine factors	
Failure of equipment/vaccine—the 36 (4)	
equipment or vaccine is faulty	
Equipment/vaccine packaging—he packaging 25 (3)	
is impractical inadequate or faulty	
Equipment/vaccine storage—inadequate 25 (3)	
impractical storage	
Poor equipment/vaccine design—the design 3 (0)	
is impractical, inadequate or faulty	
Organizational factors	
Working Conditions—factors relating to the 52 (5)	
work environment Continuity of case, issues with the	
Continuity of care—issues with the 48 (5) co-ordination of services	
Education and training—insufficient 36 (4)	
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Inadequate guidelines or protocols—existing 27 (3) guidelines not fit for purpose	
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medically and socially vulnerable children. Crosscutting reported contributory factors for quality improvement interventions include staff mistakes, coordination and verification procedures, and parents' and staff knowledge and consequent behaviors.

4.1. Strengths and limitations

This is the largest examination of reported contributory factors for immunization-related patient safety incidents in children. Incident report data are prone to numerous and well-acknowledged

biases including under-reporting and selective deposit bias [20]. For example, although the NRLS accepts reports from patients and parents, such reports were not apparent in our dataset. Further, there was likely differential reporting between organizations, those with good reporting cultures likely contribute more than those without such cultures [21].

Methodological rigor was ensured by keeping an audit trail of all coding-related decisions, holding weekly meetings to discuss analysis, and independent double-coding of 20% of reports indicating a high degree of concordance [22]. Our findings are hypothesis generating, inductive in nature, and require testing and development in further research and future clinical practice.

4.2. Reviewing these findings in the context of other literature

High numbers of reports describing administration of the wrong number of doses was expected because they are usually apparent to those involved (the healthcare professional, patient or parent) and thus more likely to be reported. Previous studies investigating immunization-related safety incidents mirror our findings and demonstrate receipt of additional vaccines is a widespread problem often resulting from poor documentation e.g. where a child is immunized but this goes undocumented resulting in that child receiving that same vaccine later [4,7,23]. In one U.S. study, over 20% of children received unnecessary duplicate vaccinations [4].

Our data contained reports of delayed vaccination or receipt of vaccines out-of-schedule with the national immunization schedule. Experts determine the vaccination schedule to afford children maximum protection from diseases and to minimize the risk of vaccine interaction. The consequences of deviating from this schedule are unclear [7,24,25].

Wrong vaccine administration is a recognized issue [7,23,26,27]. Children who receive the wrong vaccine, but in whom the error goes undetected could be both under-protected and at risk since their inadequate immunity would be unrecognized [7,23]. The potentially severe consequences of immunization-related safety incidents in medically vulnerable children have also been highlighted by other studies [28,29].

Socially vulnerable children were commonly represented in this sample and the inverse care law, where those most in need of high quality care are the least likely to receive it, is evident in this context [30]. Poorer vaccination uptake in vulnerable children is described in the literature [31–33], yet the challenges of conducting research with marginalized populations could perhaps explain the limited interventional efforts to date [34].

4.3. Recommendations for improvement

Continued efforts by manufacturers to create vaccines with different packaging and distinguishable names are needed [35,36].

Our findings support targeted community nurse visits to socially vulnerable children [37], to educate parents about the need for timely vaccination, and to encourage vaccine uptake. In addition, providing parents with access to all of their children's records could reduce documentation discrepancies and appointment-related incidents, as well as provide healthcare professionals with a safety net [38]. This could also be enhanced with better accessibility of unified immunization records for staff [39]. Building IT infrastructure and functionality capable of sharing data between health and social care providers could support identification of predictors of risk and inform interventions to mitigate future incidents [40].

Encouraging parental involvement, and creating a culture where parents feel comfortable challenging healthcare professionals, could also prevent safety incidents [38]. Co-production methods, where patients and providers co-design new models and methods of care delivery could be used to inform local improvement

initiatives that advance this parent-provider relationship for child safety [41-43]. Public health organizations and researchers must seek to establish what methods of communication work best for different patient and parent groups, and embrace the challenge of undertaking research with and for marginalized patient populations.

To reduce staff mistakes, locally-owned efforts to adopt and adapt best practices proven to be effective elsewhere should be explored for verification and standardized preparation of vaccines [36,44].

5. Conclusions

The types of immunization-related safety incidents experienced by children in England and Wales are consistent with those described in studies undertaken in other countries. This is the largest examination of reported contributory factors for immunization-related patient safety incidents in children, and based on this analysis we have made a number of priority recommendations for policy, practice, and research. These include: investments in IT infrastructure to support data linkage and the identification of risk predictors; development of consultation models to enhance parental roles in mitigating safety incidents; and improvement efforts to adapt and adopt best practices from elsewhere. These priorities are particularly pressing for medically and socially vulnerable patients.

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Contributor statements

Ms Rees was responsible for primary data coding, qualitative analysis, interpretation of results, and drafting the manuscript. Professor Edwards conceptualized and designed the study. He carried out interpretation of the results, created the recommendations for practice and reviewed the manuscript. Dr Powell provided the clinical interpretation of results, creation of recommendations for pediatric practice, and reviewed the manuscript. Dr Carter was the study statistician and responsible for data analysis, and interpretation of the results and reviewed the manuscript. Dr Evans was the project clinical informatics lead and responsible for development of the data storage infrastructure, clinical interpretation of results and creating recommendations for improvement. Mr Hibbert conceptualized and designed the study and development of the in-house classification systems. He interpreted the results and reviewed the manuscript. Dr Makeham contributed to the clinical interpretation of results, creation of recommendations for family practice, and reviewed the manuscript. Professor Sheikh conceptualized and designed the study interpretation of results, and reviewed the manuscript. Professor Donaldson conceptualized and designed the study interpretation of results, and reviewed the manuscript. Dr Andrew Carson-Stevens conceptualized and designed the study. He was responsible for data coding, qualitative analysis, interpreting results, developing recommendations for clinical practice, and revising the manuscript. He is also the study guarantor. All co-authors approved the final manuscript as submitted.

Conflict of interest statement

ACS and AE are co-chief investigators, and AS and LD are coapplicants on a NIHR HS&DR funded study to characterize primary care patient safety incident reports.

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Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at http://dx.doi.org/10.1016/j.vaccine.2015.06.

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Appendix 2

2.1 Search terms used to retrieve vaccination-related reports

- 1. BCG
- 2. Booster
- 3. Cervarix
- Conjugate
 Diphtheria
 DTP

- 7. DTaP 8. Gardasil 9. Havrix
- 10. Haemophilus
- 11. Hep A
- 12. Hep B
- 13. Hepatitis A
- 14. Hib
- 15. HPV
- 16. Human papilloma
- 17. Imms
- 18. Immuni
- 19. Infanrix
- 20. Influenza
- 21. Inject
- 22. IPV
- 23. Jab
- 24. Measles
- 25. Men c
- 26. MenC
- 27. Meningitis C
- 28. Meningococcal
- 29. Menitorix
- 30. MMR
- 31. Mumps
- 32. Pandemrix
- 33. PCV
- 34. Pediacel
- 35. Pertussis
- 36. Pneumococcal
- 37. Polio
- 38. Prevenar
- 39. Priorix
- 40. PSB
- 41. Red book
- 42. Repevax
- 43. Revaxis
- 44. Rotavirus
- 45. Rubella
- 46. Tetanus
- 47. Td/IPV
- 48. Typhoid 49. Vaccin
- 50. Vacs
- 51. Varicella

2.2 Search terms used to retrieve reports involving 'unwell' children

- 1. Abdo pain
- 2. Abdominal pain
- 3. Acetazolamide
- 4. Acute abdo
- 5. Anorexi
- 6. Antibiot
- 7. Apnoe
- 8. Appendicitis
- 9. Arrest
- 10. Arthritis
- 11. ASD
- 12. Asperg
- 13. Asthma
- 14. Ataxia
- 15. Atomoxetine
- 16. Atresia
- 17. Autis
- 18. Bacteria
- 19. Behavioural
- 20. Blanch
- 21. Blastoma
- 22. Blind
- 23. Blood sugar
- 24. Breathless
- 25. Bronchiolitis
- 26. Bronchitis
- 27. Burn
- 28. CAMHS
- 29. Cancer
- 30. Carbamazepine
- 31. Cardiac failure
- 32. Cellulitis
- 33. Cerebral palsy
- 34. Chemo
- 35. Citalopram
- 36. Clobazam
- 37. Clomipramine
- 38. Clonazepam
- 39. Clonidine
- 40. CMHT
- 41. Coarctation of the Aorta
- 42. Coelia disease
- 43. Coeliac disease
- 44. Coma
- 45. Congenital
- 46. Convulsion
- 47. Cough
- 48. Cramp
- 49. Crohn
- 50. Cyanos
- 51. Cystic
- 52. Deaf
- 53. Dehydrate
- 54. Depress

- 55. Deteriorate
- 56. Development delay
- 57. Diabet
- 58. Dialys
- 59. Diarrhea
- 60. Diarrhoea
- 61. Diazepam
- 62. Dipstick
- 63. Disab
- 64. Disorder
- 65. Dissociative
- 66. DKA
- 67. Down syndrome
- 68. Downs syndrome
- 69. Drowsy
- 70. Duloxetine
- 71. Dystrophy
- 72. E.coli
- 73. Ectomy
- 74. Eczema
- 75. Ehlers Danlos
- 76. Epilep
- 77. Erythema multiforme
- 78. Ethosuximide
- 79. Exacerbate
- 80. Eve syndrome
- 81. Foetal alcohol
- 82. Febrile
- 83. Fever
- 84. Floppy
- 85. Fluoxetine,
- 86. Fracture
- 87. Gabapentin
- 88. Gastritis
- 89. Gastritis
- 90. Gastroenteritis
- 91. GCS
- 92. George syndrome
- 93. Glucose
- 94. Glycosuria
- 95. Growth hormone
- 96. Haematuria
- 97. Haemol
- 98. Haemophilia
- 99. Haloperidol
- 100. Head injury
- 101. Headache
- 102. Heart
- 103. Hirschprung syndrome
- 104. HIV
- 105. Hydrocephalus
- 106. Hyper
- 107. Hypo
- 108. Imipramine
- 109. Immuno
- 110. Impetigo
- 111. Infect
- 112. Inhaler
- 113. Insulin
- 114. Intra osseous

- 115. Intra venous
- 116. Intracranial
- 117. Intubate
- 118. Intussusception
- 119. Irritable
- 120. IUGR
- 121. IV access
- 122. IV antihistamine
- 123. IV diazepam
- 124. IV fluid
- 125. IV sedation
- 126. Jaundice
- 127. Kawazaki disese
- 128. Ketoacidosis
- 129. Keton
- 130. Ketosis
- 131. Kidney
- 132. Lamotrigine
- 133. Learning disa
- 134. Leukae
- 135. Levetiracetam
- 136. Life saving
- 137. Life threatening
- 138. Life-saving
- 139. Lithium
- 140. Liver disease
- 141. Liver failure
- 142. Metab
- 143. Acido
- 144. Renal failure
- 145. Phenobarbitone
- 146. Sepsis
- 147. Lorazepam
- 148. lung disease
- 149. Lung disease
- 150. Lymph
- 151. Maln
- 152. Meningitis
- 153. Meningitus
- 154. Meningoco
- 155. Meningococcal disease
- 156. Methylphenidate
- 157. Midazolam
- 158. Migraine
- 159. Mirtazapine
- 160. Myopathy
- 161. Nebs
- 162. Nebu
- 163. Nephrotic syndrome
- 164. Neuroblastoma
- 165. Nitrazepam
- 166. Obstruct
- 167. Oedema
- 168. Olanzapine
- 169. Osteomyelitis
- 170. Otitis
- 171. Otitis Media
- 172. Overdose
- 173. Oxcarbazepine
- 174. Palsy

- 175. Paroxetine
- 176. Pendred syndrome
- 177. Perthes disease
- 178. Petech
- 179. Phenobarbital
- 180. Phenytoin
- 181. Plegia
- 182. Pneumococcal disease
- 183. Pneumonia
- 184. Pneumonitis
- 185. Polyuria
- 186. Promazine
- 187. Psychiat
- 188. Psychol
- 189. Psychosis
- 190. Pulse
- 191. Pyrexia
- 192. Quetiapine
- 193. Qunisy
- 194. Rash
- 195. Refeeding syndrome
- 196. Reflux
- 197. Renal disease
- 198. Renal failure
- 199. Renal func
- 200. Resp rate
- 201. Respiratory rate
- 202. Responsive
- 203. Retts syndrome
- 204. Rey's syndrome
- 205. Risperidone
- 206. Ritalin
- 207. Salbutamol
- 208. Sats
- 209. Saturation
- 210. Scabies
- 211. Scalds
- 212. Scoliosis
- 213. Seizure
- 214. Self harm
- 215. Self-harm
- 216. Septic
- 217. Septic
- 218. Croup
- 219. Obstruct
- 220. Oedema
- 221. Sertraline
- 222. Shortness of breath
- 223. Sick
- 224. Sickle cell disease
- 225. Sickle disease
- 226. Sodium valproate
- 227. Spina bifida
- 228. Spinal muscular atrophy
- 229. Splenectomy
- 230. Squint
- 231. Staph
- 232. Steroid
- 233. Strattera
- 234. Streptococcal

- 235. Suicide
- 236. SVT
- 237. Swollen
- 238. Tachy
- 239. Talipes
- 240. Tender
- 241. Thalassaemia
- 242. Thrive
- 243. Thrombo
- 244. Thyroid
- 245. Tonsillitis
- 246. Topiramate
- 247. Torticollis
- 248. Transposition of the great arteries
- 249. Tuberous sclerosis
- 250. Tumour
- 251. Undescended
- 252. Urinary tract
- 253. Urticaria
- 254. Valproate
- 255. Vasculitis
- 256. Ventilation
- 257. Ventolin
- 258. Viral
- 259. Visual impairment
- 260. Von willebrand
- 261. VSD
- 262. Westerdrout syndrome
- 263. Wheeze
- 264. Zolpidem

2.3 Incident descriptors framework

1 ** ADMINISTRATION **

- 1.1 Filing system information filed incorrectly
- 1.2 Message handling errors in the taking and distributing of messages
- 1.3 Appointments errors in managing appointments for healthcare
 - 1.3.1 Primary care appointments
 - 1.3.2 Secondary care appointments
- 1.4 Payment errors in the process of healthcare payment systems
- 1.5 Ability to access healthcare professional
 - 1.5.1 Home visits professional delayed/ unable to visit patient at home
 - 1.5.2 Returning phone calls
 - 1.5.3 Out-of-hours
 - 1.5.4 Health visiting
 - 1.5.5 Child and Adolescent Mental Health Services
 - 1.5.6 Occupational therapy
- ${\bf 1.6\ Transfer\ of\ patient\ information\ -\ incorrect\ or\ inefficient\ transfer\ of\ patient\ information\ across\ healthcare\ systems}$
 - 1.6.1 Between care settings
 - 1.6.1.1 From primary to secondary care
 - 1.6.1.1.1 Lost
 - 1.6.1.1.2 Not sent
 - 1.6.1.1.3 Incorrect/incomplete
 - 1.6.1.1.4 Delayed
 - 1.6.1.1.5 Illegible
 - 1.6.1.2 From secondary to primary care
 - 1.6.1.2.1 Lost
 - 1.6.1.2.2 Not sent
 - 1.6.1.2.3 Incorrect/incomplete
 - 1.6.1.2.4 Delayed
 - 1.6.1.2.5 Illegible
 - 1.6.1.3 Between primary care settings
 - 1.6.1.3.1 Lost
 - 1.6.1.3.2 Not sent
 - 1.6.1.3.3 Incorrect/incomplete
 - 1.6.1.3.4 Delayed
 - 1.6.1.3.5 Illegible
 - 1.6.2 New diagnoses incorrect or inefficient transfer of patient information from secondary care regarding new diagnoses

- 1.6.3 Appropriate follow up incorrect or inefficient transfer of patient regarding necessary follow-up of patient. e.g. requirements for follow up screening or regular review
- 1.6.4 Involving out-of-hours incorrect or inefficient transfer of patient information between in- and out- of hours services
- 1.6.5 NHS direct
- 1.7 Breaches of confidentiality

2 ** DOCUMENTATION**

- 2.1 Medical records
 - 2.1.1 Record(s) unavailable records could not be accessed when needed
 - 2.1.1.1 Red book
 - 2.1.1.2 General practice records
 - 2.1.1.3 Child health records
 - 2.1.1.4 Lost medical records
 - 2.1.2 Care given but not documented
 - 2.1.3 Record not up to date or complete information missing from records
 - 2.1.3.1 Discrepancies between vaccine records
 - 2.1.3.1.1 Red book
 - 2.1.3.1.2 General practice records
 - 2.1.3.1.3 Child health records
 - 2.1.4 Inaccurate or unclear medical records / medical record error
 - 2.1.4.1 Red book
 - 2.1.4.2 General practice records
 - 2.1.4.3 Child health records
- 2.2 Death certificates

3 ** REFERRAL **

- 3.1 Human
 - 3.1.1 Not performed when indicated
 - 3.1.1.1 Delayed referral errors in the timely referral of patients
 - 3.1.1.1 Secondary care
 - 3.1.1.1.2 Specialist care
 - 3.1.1.3 Emergency care
 - 3.1.1.1.4 Nursing
 - 3.1.1.1.5 Social care

- 3.1.1.1.6 Health visitor
- 3.1.1.7 General practice
- 3.1.1.1.8 Child and Adolescent Mental Health Services
- 3.1.1.2 Referral not made when appropriate referral decision-making error
 - 3.1.1.2.1 Secondary care
 - 3.1.1.2.2 Specialist Care
 - 3.1.1.2.3 Emergency Care
 - 3.1.1.2.4 Nursing
 - 3.1.1.2.5 Health visitor
 - 3.1.1.2.6 Social care
 - 3.1.1.2.7 General practice
 - 3.1.1.2.8 Child and Adolescent Mental Health Service
- 3.1.1.3 No follow up arranged did not follow-up or were not asked to follow-up
- 3.1.2 Incomplete /incorrect referral
- 3.1.3 Illegible referral
- 3.1.4 Work inappropriately passed to primary care
- 3.1.5 Inappropriate referral
- 3.1.6 Referral refused

3.2 Administration

- 3.2.1 Not sent- letter of referral erroneously not sent by office
- 3.2.2 Delayed letter of referral delayed at office level
- 3.2.3 Lost letter of referral lost in the system
- 3.2.4 Not acted upon referral successful but patient not seen by physic 3.2.4.1 Refused patient referral refused by receiving office
- 3.2.5 Inappropriate referral-referral made erroneously at office level
- 3.2.6 Social work referral issues

4 ** DIAGNOSIS AND ASSESSMENT **

- 4.1 Diagnosis
 - 4.1.1 Missed diagnosis
 - 4.1.2 Wrong diagnosis
 - 4.1.3 Delayed diagnosis
 - 4.1.3.1 Cancer
 - 4.1.3.2 Emergency condition
 - 4.1.3.3 Contagious condition
- 4.2 Insufficient assessment

- 4.2.1 Triage errors in the process of triaging patients
 - 4.2.1.1 By healthcare professional
 - 4.2.1.2 By non-healthcare professional
- 4.2.2 History errors in the process of taking a patient's medical history
- 4.2.3 Examination errors in the process of examining patients
- 4.2.4 Identifying vulnerable or high-risk patient
- 4.2.5 Emergency vehicle use Inappropriate transfer vehicle used (e.g. private vehicle instead of ambulance)
- 4.2.6 Discharge planning premature discharge and poor discharge planning
- 4.3 Delayed assessment a delay in assessment for care or care adjunct

5 ** TREATMENT & PROCEDURES (excludes drugs/vaccines)

- 5.1 Clinical treatment decision
 - 5.1.1 No treatment given
 - 5.1.2 Insufficient treatment given
 - 5.1.3 Wrong treatment given
- 5.2 Treatment other than medication
 - 5.2.1 Ordering treatments wrong treatment ordered or treatment not ordered when appropriate
 - 5.2.2 Implementation error in conducting the correctly chosen process or procedure
 - 5.2.3 Complication
 - 5.2.3.1 Complication from execution of procedure
 - 5.2.3.2 Adverse event suffered by patient as a result of treatment other than medication
 - 5.2.4 Timeliness treatment other than medication not administered in a timely fashion
 - 5.2.5 Execution of care error in choosing the correct process or procedure
 - 5.2.6 Wrong anatomical side/site
 - 5.2.7 Insufficient supply of treatment

6 ** MEDICATION & VACCINES **

- 6.1 Clinical treatment decision
 - 6.1.1 No treatment given

- 6.1.2 Insufficient treatment
- 6.1.3 Wrong treatment given
- 6.1.4 Treatment not ordered

6.2 Medication prescribing

- 6.2.1 Wrong medication
- 6.2.2 Wrong patient
- 6.2.3 Wrong dose
- 6.2.4 Wrong route
- 6.2.5 Wrong time
- 6.2.6 Unsafe medication
 - 6.2.6.1 Teratogenic
 - 6.2.6.2 Contraindicated
 - 6.2.6.3 Allergy prescribed for patient with known allergy
- 6.2.7 Wrong formulation
- 6.2.8 Wrong number of doses
- 6.2.9 Illegible/ unclear prescription
- 6.2.10 Incomplete prescription e.g. brand not specified

6.3 Medication dispensing

- 6.3.1 Wrong medication
- 6.3.2 Wrong patient
- 6.3.3 Wrong dose
- 6.3.4 Wrong route
- 6.3.5 Wrong time
- 6.3.6 Wrong formulation
- 6.3.7 Not dispensed
- 6.3.8 Allergy dispensed to a patient with known allergy
- 6.3.9 Out of date
- 6.3.10 Wrong label
- 6.3.11 Wrong number of doses
- 6.3.12 Inappropriate medication
- 6.3.13 Medication dispensed in inappropriate container

6.4 Medication administration

- 6.4.1 Wrong medication
- 6.4.2 Wrong patient
- 6.4.3 Wrong dose
- 6.4.4 Wrong route
- 6.4.5 Wrong time
- 6.4.6 Wrong formulation
- 6.4.7 Out of date
- 6.4.8 Allergy medication administered to patient with known allergy
- 6.4.9 Medication not administered
- 6.4.10 Reconstitution error

6.5 Monitoring medication - error in the process of monitoring dose-dependent medications, or those with side effects

- 6.5.1 Lack of monitoring
- 6.5.2 Medication dose not appropriately adjusted

- 6.6 Adverse event patient suffered a complication as a result of medication
 - 6.6.1 Allergy unknown that patient had any allergies
- 6.7 Drug omission medication erroneously not given to or not taken by patient
- 6.8 Patient self-administered overdose
- 6.9 Incorrect storage
- 6.10 Medication timeliness not commenced in a timely fashion

6.11 Vaccines

6.11.1 Vaccine prescribing

- 6.11.1.1 Wrong vaccine
- 6.11.1.2 Wrong patient
- 6.11.1.3 Wrong dose
- 6.11.1.4 Wrong route
- 6.11.1.5 Wrong time
- 6.11.1.6 Contraindicated
- 6.11.1.7 Wrong formulation
- 6.11.1.8 Wrong number of doses

6.11.2 Vaccine dispensing

- 6.11.2.1 Wrong vaccine
- 6.11.2.2 Wrong patient
- 6.11.2.3 Wrong dose
- 6.11.2.4 Wrong route
- 6.11.2.5 Wrong time
- 6.11.2.6 Wrong number of doses
- 6.11.2.7 Stored incorrectly
- 6.11.2.8 Out of date
- 6.11.2.9 Not dispensed
- 6.11.2.10 Wrong formulation
- 6.11.2.11 Wrong label
- 6.11.2.12 Contraindicated

6.11.3 Vaccine administration

- 6.11.3.1 Wrong vaccine
- 6.11.3.2 Wrong patient
- 6.11.3.3 Wrong dose
- 6.11.3.4 Wrong route
- 6.11.3.5 Wrong time
- 6.11.3.6 Wrong amount
- 6.11.3.7 Stored incorrectly
- 6.11.3.8 Out of date
- 6.11.3.9 Contraindicated vaccine
- 6.11.3.10 Not administered
- 6.11.3.11 Used/dirty needle
- 6.11.3.12 Wrong site
- 6.11.4 Reconstitution error
- 6.11.5 Adverse event (reaction to vaccine)
- 6.11.6 Batch recall

6.12 Vaccine unavailable

7 ** INVESTIGATIONS **

- 7.1 Laboratory errors in the process of laboratory investigations
 - 7.1.1 Ordering wrong test ordered or test not ordered when appropriate 7.1.2 Implementing errors in the process of obtaining or processing a laboratory specimen
 - 7.1.2.1 Mislabelled sample
 - 7.1.3 Reporting error in the process of physician receiving accurate test results including errors of delay
 - 7.1.4 Responding to results inappropriate response to a laboratory result
- 7.2 Diagnostic imaging errors in the process of diagnostic imaging investigations
 - 7.2.1 Ordering wrong test ordered or test not ordered when appropriate 7.2.2 Implementing errors in the process of obtaining or processing of a diagnostic image
 - 7.2.2.1 Mislabelled request form
 - 7.2.3 Reporting error in the process of physician receiving accurate test results including errors of delay
 - 7.2.4 Responding to results inappropriate response to a laboratory result
- 7.3 Other investigations
 - 7.3.1 Ordering wrong test ordered or test not ordered when appropriate
 - 7.3.2 Implementing errors in the process of obtaining or processing of other diagnostic investigation
 - 7.3.3 Reporting error in the process of physician receiving accurate test results including errors of delay
 - $7.3.4 \ Responding$ to results inappropriate response to a result of other investigations

8 ** COMMUNICATION **

These are human failures, and do not include breakdowns in the systems that are used to communicate information.

- 8.1 With patients or caregivers errors in communication between physicians or healthcare professionals and patients or caregivers
 - 8.1.1 Wrong advice given to patient or caregiver includes information about accessing emergency services, self-management or safety netting
 - 8.1.1.1 By healthcare professional
 - 8.1.1.2 By non-healthcare professional
 - 8.1.2 Failure to convey seriousness/urgency of patient condition
 - 8.1.3 Consent errors errors in the process of obtaining informed consent
- 8.2 Between healthcare professionals errors in communication between healthcare professionals

- 8.2.1 Failure to convey seriousness/urgency of patient condition
- 8.2.2 Handover-related inadequacies
- 8.3 Between healthcare and non-healthcare professionals

9 ** EQUIPMENT **

- 9.1 Therapeutic adjunct provision
- 9.2 Insufficient supply
- 9.3 Failure of equipment
 - 9.3.1 Damaged
 - 9.3.2 Faulty
 - 9.3.3 Misused
 - 9.3.4 Computerised Physician Order Entry
- 9.4 Stolen equipment

10 ** OTHER **

- 10.1 Professionalism
- 10.2 Environmental hazard
- 10.3 Transport issues
- 10.4 Failure to prevent fall/injury
- 10.5 Failure to follow up 'unwell' or vulnerable child
- 10.6 Failure to prevent pressure ulcer

2.4 Contributory factors framework

1 ** PATIENT OR CAREGIVER FACTORS **

- 1.1 Geography the area where patients live including its characteristics
 - 1.1.1 Out of area patient new to area
 - 1.1.2 Access difficulties because of geography
- 1.2 Language patient unable to communicate in English
- 1.3 Behaviour
 - 1.3.1 Non-compliance patient does not follow advice or instructions
 - 1.3.1.1 Takes own discharge
 - 1.3.1.2 Patient does not take medication as instructed or advise
 - 1.3.1.3 Non-disclosure
 - 1.3.1.4 Violent
- 1.4 Health factors related to the patient's physical and mental health
 - 1.4.1. Frailty reduced physiological reserve, fragile
 - 1.4.2. Disability
 - 1.4.3. Allergy
 - 1.4.4 Immunocompromised
 - 1.4.5 Coagulation problems
 - 1.4.6 Pregnancy
 - 1.4.7 Epilepsy
 - 1.4.8 Poor renal function
- 1.5. Knowledge patient or caregiver of child has poor understanding
- 1.6. Looked-after child
- 1.7 Age
 - 1.7.1 Weight-based dosing
- 1.8 Ethnicity

2 ** STAFF FACTORS **

- 2.1 Physical and mental wellbeing
 - 2.1.1 Fatigue
- 2.2 Task a piece of work to be done or undertaken.
 - 2.2.1 Failure to follow protocol
 - 2.2.1.1 New protocol
 - 2.2.2 Inadequate skill set/knowledge

- 2.3 Cognitive includes abilities such as perception, learning, memory, language, concept formation, problem solving, and thinking.
 - 2.3.1 Mistake
 - 2.3.1.1 Distraction/ inattention/ oversight/forgot
 - 2.3.1.2 Similar medication names / appearances confused
 - 2.3.1.3 Similar patient names
 - 2.3.1.4 Haste/ poor time management
 - 2.3.1.5 Misread/ did not read
 - 2.3.1.6 Wrong patient's sticky label
 - 2.3.2 Violation deliberate breaking of a rule
 - 2.3.3 Stress mental or emotional strain
 - 2.3.4 No or poor supervision or assistance of staff

3 ** EQUIPMENT / MEDICATION/ VACCINE FACTORS **

- 3.1 Poor design impractical, faulty or in some way inadequate
- 3.2 Poor storage
- 3.3 Poor packaging
- 3.4 Failure of equipment/ medication/ vaccine

4 ** ORGANISATION FACTORS **

- 4.1 Protocols or guidelines inadequate, inefficient absent or not available
 - 4.1.1 Mental health
 - 4.1.2 Vulnerable patients
 - 4.1.3 Investigations
 - 4.1.4 Referrals
 - 4.1.5 Epilepsy management plan
 - 4.1.6 Asthma management plan
 - 4.1.7 School care plan
 - 4.1.8 Diabetic management plan
 - 4.1.9 Palliative care plan
- 4.2 Interpreter services communication aids to reduce language barriers
- 4.3 Continuity of care the delivery of a 'seamless service' through integration, coordination and the sharing of information between different providers
 - 4.3.1 Unknown to staff
 - 4.3.2 Within primary care
 - 4.3.2.1 Out-of-hours service

4.3.2.2 Registering with a general practice

- 4.3.3 Between secondary and primary care
- 4.3.4 Access block cannot move a patient on because there is no space to put them
- 4.3.5 Locum/ agency staff

4.4 Working conditions

- 4.4.1 Staffing levels
 - 4.4.1.1 Shift pattern
 - 4.4.1.2 Insufficient numbers of staff
 - 4.4.1.2.1 Doctors
 - 4.4.1.2.2 Nurses
 - 4.4.1.2.3 Allied health professionals
 - 4.4.1.3. Sickness
- 4.4.2 Team factors
 - 4.4.2.1 Culture
 - 4.4.2.2 Inadequate leadership
 - 4.4.2.3 Disagreement amongst teams
- 4.4.3 Busy/overloaded by work
- 4.4.4 interruptions
- 4.5. Education and training
 - 4.5.1 Supervision
 - 4.5.2 Knowledge of others roles
 - 4.5.3 Caregiver training
- 4.6 Service unavailable
- 4.7 Long wait for service

5 ** ENVIRONMENTAL FACTORS **

5.1 Care facility has poor access for emergency vehicles

2.5 Incident outcomes framework

1 ** PATIENT HARM **

1.1 Clinical harm

- 1.1.1 Pain / discomfort
- 1.1.2 Swelling
- 1.1.3 Rash
- 1.1.4 Nausea
- 1.1.5 Redness
- 1.1.6 Bruising
- 1.1.7 Dizziness/ faint/ loss or altered consciousness
- 1.1.8 Bleeding
- 1.1.9 Changes in physiological parameters
 - 1.1.9.1 Fever
 - 1.1.9.2 Breathless
- 1.1.10 General deterioration/progression of condition
- 1.1.11 Pressure ulcer
 - 1.1.11.1 Pressure ulcer developed
 - 1.1.11.2 Pressure ulcer deteriorated
- 1.1.12 Other wound/ulcer
- 1.1.13 Admitted to the high dependency or intensive care unit
- 1.1.14 Seizures
- 1.1.15 Admitted to hospital/visited emergency department
- 1.1.16 Infection
- 1.1.17 Migraine
- 1.1.18 Poor diabetic control
 - 1.1.18.1 Diabetic ketosis/ ketoacidosis
- 1.1.19 Developmental delay
- 1.1.20 Diarrhoea
- 1.1.21 Emergency surgery
- 1.1.22 Liver failure
- 1.1.23 Constipation

1.2 Injury

- 1.2.1 Laceration
- 1.2.2 Perforation
- 1.2.3 Fracture
- 1.2.4 Skin tear
- 1.2.5 Pain / discomfort
- 1.2.6 Swelling
- 1.2.7 Redness
- 1.2.8 Bruising
- 1.2.9 Bleeding
- 1.2.10 Needle stick
- 1.2.11 Burn
- 1.2.12 Fall

1.3 Psychological / emotional distress

- 1.4 Death
- 1.5 Cardio-respiratory arrest

2 ** PATIENT INCONVENIENCE **

- 2.1 Repeated tests / procedure / additional treatment
- 2.2 Delays in management (assessment or treatment)
- 2.3 Increased documentation
- 2.4 Financial implication
- 2.5 Repeated visits to/from health care providers
- 2.6 Unnecessary treatment
- 2.7 Extended hospital stay
- 2.8 Hospital admission

3 ** ORGANISATIONAL INCONVENIENCE **

- 3.1 Increased documentation
- 3.2 Phone calls/follow-up
- 3.3 More equipment / supplies used
- 3.4 Delays in using facilities
- 3.5 Legal implication
- 3.6 Complaint made
- 3.7 Financial implication

Appendix 3

3.1 The frequencies of combinations of incident types described by included vaccination-related reports

	Combinations of incident types	Fr	requency of
		C	ombination
Wrong vaccine			218
Wrong number of			173
doses			
Wrong timing			145
Adverse reaction			140
Wrong number of	Out of date		76
doses	records		
Procedural error			57
Out of date			54
records			
Not administered	Reconstitution		50
	error		
Expired vaccine			46
administered			
Wrong number of	Wrong vaccine		46
doses			
Wrong timing	Appointment		45
	management		
Communication -			43
patients/			
caregivers			
Wrong number of	Record		31
doses	availability		
Appointment			29
management			
Records			29
inaccurate/			
unclear			
Vaccine			24
prescribing and			
dispensing			

incident			
Wrong dose			24
Wrong number of	Out of date	Record	23
doses	records	availability	
Wrong dose	Procedural error		23
Wrong number of	Communication -		19
doses	patients/		
	caregivers		
Wrong patient			16
Wrong timing	Communication -		16
	patients/		
	caregivers		
Transfer of			15
information			
Vaccine not			15
administered			
Non-specific			14
vaccine			
administration			
incident			
Wrong timing	Transfer of		13
	information		
Wrong number of	Appointment		12
doses	management		
Wrong number of	Records		12
doses	inaccurate/		
	unclear		
Contraindicated			10
vaccine			
Wrong vaccine	Record		9
	availability		
Wrong timing	Record		9
	availability		
Wrong timing	Wrong vaccine		9
Wrong number of	Out of date		8
doses	records		
Wrong number of	Out of date	Communication -	7
doses	records	patients/	
		caregivers	

Wrong number of	Transfer of		7
doses	information		
Wrong timing	Records		7
	inaccurate/		
	unclear		
Vaccine			6
administered at			
wrong site			
Wrong number of	Wrong patient		6
doses			
Wrong vaccine	Wrong patient		6
Wrong timing	Appointment	Transfer of	6
	management	information	
Appointment	Out of date		5
management	records		
Environmental			5
hazard			
Record			5
availability			
Wrong vaccine	Records		5
	inaccurate/		
	unclear		
Wrong dose	Vaccine		5
	prescribing and		
	dispensing		
	incident		
Wrong timing	Out of date		5
	records		
Appointment	Transfer of		4
management	information		
Batch recall			4
Reconstitution			4
error			
Out of date	Transfer of		4
records	information		
Wrong number of	Appointment	Out of date	4
doses	management	records	
Wrong number of	Record	Communication -	4
doses	availability	patients/	

		caregivers	
Wrong number of	Record	Out of date	4
doses	availability	records	
Wrong number of	Out of date	Transfer of	4
doses	records	information	
Adverse reaction	Communication -		3
	patients/		
	caregivers		
Communication -	Out of date		3
patients/	records		
caregivers			
Out of date	Record		3
records	availability		
Out of date	Out of date		3
records	records		
Referral for			3
vaccination			
Used needle			3
Vaccine			3
administered via			
wrong route			
Vaccine not	Appointment		3
administered	management		
Vaccine not	Procedural error		3
administered			
Vaccine not	Reconstitution	Insufficient	3
administered	error	vaccine supply	
Vaccine stored			3
incorrectly			
Wrong number of	Record		3
doses	availability		
Wrong timing	Appointment	Non-specific	3
	management	administrative	
	J	incident	
Wrong timing	Out of date	Record	3
	records	availability	
Communication -		,	2
professionals			
Communication -	Record		2
3.000			_

Caregivers Communication - patients / Caregivers Contraindicated	patients/	availability		
patients/ caregivers Contraindicated vaccine professionals Contraindicated vaccine patients/ caregivers Contraindicated vaccine patients/ caregivers Contraindicated vaccine inaccurate/ unclear Non-specific vaccine administration incident Other administrative incident Used needle Procedural error vaccine not administered Vaccine appointment prescribing and dispensing incident Wrong number of doses professionals Wrong number of Insufficient vaccine supply Wrong number of Justificient vaccine supply	caregivers			
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Wrong number of Out of date Communication - 2	Wrong number of	Insufficient		2
	doses	vaccine supply		
doses records patients/	Wrong number of	Out of date	Communication -	2
,	doses	records	patients/	

		caregivers	
Wrong number of	Out of date	Record	2
doses	records	availability	
Wrong number of	Transfer of	Record	2
doses	information	availability	
Wrong vaccine	Communication -		2
_	patients/		
	caregivers		
Wrong vaccine	Out of date	Record	2
	records	availability	
Wrong dose	Wrong vaccine		2
Wrong timing	Appointment	Communication -	2
	management	patients/	
		caregivers	
Wrong timing	Appointment	Records	2
	management	inaccurate/	
		unclear	
Wrong timing	Appointment	Out of date	2
	management	records	
Wrong timing	Communication -		2
	professionals		
Wrong timing	Non-specific		2
	documentation		
	incident		
Wrong timing	Other		2
	administrative		
	incidents		
Wrong timing	Records	Appointment	2
	inaccurate/	management	_
	unclear	, manuage mene	
Adverse reaction	Appointment		1
	management		
Adverse reaction	Contraindicated	Communication -	1
	vaccine	patients/	
		caregivers	
Adverse reaction	Out of date		1
	records		
Appointment	Communication -		1
management	professionals		
	F		

vaccine error with patients/ unclear Contraindicated vaccine assessment patients/ incident caregivers Contraindicated Investigation incident vaccine records Insufficient vaccine supply Medication error Medication error Adverse reaction Non-specific vaccine patients/ administration incident Non-specific vaccine availability records administration incident Non-specific vaccine records administration incident records	Communication -	Appointment	Record	1
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vaccine records Insufficient vaccine supply Medication error Medication error Adverse reaction Non-specific Communication - vaccine patients/ administration caregivers incident Non-specific Record vaccine availability administration incident Non-specific Record Communication - vaccine availability patients/ caregivers Non-specific vaccine availability caregivers Non-specific Vaccine availability patients/ administration incident Non-specific Out of date vaccine records administration administration administration	vaccine	incident		
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vaccine availability patients/ administration incident Non-specific Out of date vaccine administration vaccine administration vaccine administration	incident			
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incident Non-specific Out of date vaccine records administration	vaccine	availability	patients/	
Non-specific Out of date vaccine records administration	administration		caregivers	
vaccine records administration	incident			
administration	Non-specific	Out of date		1
	vaccine	records		
	administration			
incident	incident			
Non-specific Referral for Records	Non-specific	Referral for	Records	1
vaccine vaccination inaccurate/	vaccine	vaccination	inaccurate/	
administration unclear	administration		unclear	
incident	incident			
Non-specific	Non-specific			1
vaccination-	vaccination-			

related incident			
Out of date	Insufficient		1
vaccine	vaccine supply		
administered			
Out of date	Vaccine		1
vaccine	prescribing and		
administered	dispensing		
	incident		
Out of date	Wrong vaccine		1
vaccine			
administered			
Professionalism			1
Professionalism	Communication -		1
	patients/		
	caregivers		
Record	Out of date		1
availability	records		
Records	Out of date		1
inaccurate/	records		
unclear			
Records	Referral for	Out of date	1
inaccurate/	vaccination	records	
unclear			
Out of date	Communication -		1
records	professionals		
Out of date	Communication -		1
records	patients/		
	caregivers		
Transport	Adverse reaction		1
Vaccine not	Communication -	Procedural error	1
administered	patients/		
	caregivers		
Vaccine not	Insufficient		1
administered	vaccine supply		
Vaccine not	Record		1
administered	availability		
Vaccine not	Wrong vaccine		1
administered			
Vaccine stored	Communication -		1

incorrectly	professionals			
Wrong number of	Appointment	Records		1
doses	management	inaccurate/		
		unclear		
Wrong number of	Communication -	Record		1
doses	professionals	availability		
Wrong number of	Communication -	Out of date		1
doses	professionals	records		
Wrong number of	Communication -	Appointment	Out of date	1
doses	patients/	management	records	
	caregivers			
Wrong number of	Communication -	Out of date		1
doses	patients/	records		
	caregivers			
Wrong number of	Communication -	Transfer of		1
doses	patients/	information		
	caregivers			
Wrong number of	Other			1
doses	administrative			
	incidents			
Wrong number of	Record	Communication -		1
doses	availability	professionals		
Wrong number of	Record	Non-specific		1
doses	availability	administrative		
		incident		
Wrong number of	Record	Transfer of		1
doses	availability	information		
Wrong number of	Records	Appointment	Out of date	1
doses	inaccurate/	management	records	
	unclear			
Wrong number of	Records	Record		2
doses	inaccurate/	availability		
	unclear			
Wrong number of	Out of date	Communication -		1
doses	records	professionals		
Wrong number of	Out of date	Non-specific	Record	1
doses	records	administrative	availability	
		incident		
Wrong number of	Out of date	Record	Communication -	1
	<u> </u>			

doses	records	availability	patients/	
			caregivers	
Wrong number of	Out of date	Records		1
doses	records	inaccurate/		
		unclear		
Wrong number of	Out of date	Records	Communication -	1
doses	records	inaccurate/	patients/	
		unclear	caregivers	
Wrong number of	Out of date		ou. og. v o. o	1
doses	records			
Wrong number of	Transfer of	Communication -		1
doses	information	patients/		
doses	in ormación	caregivers		
Wrong number of	Transfer of	Out of date		1
	information			1
doses		records		
Wrong number of	Wrong vaccine	Record		1
doses		availability		
Wrong number of	Wrong vaccine	Record	Appointment	1
doses		availability	management	
Wrong number of	Wrong vaccine	Records		1
doses		inaccurate/		
		unclear		
Wrong number of	Wrong vaccine	Wrong patient		1
doses				
Wrong vaccine	Appointment			1
	management			
Wrong vaccine	Insufficient			1
	vaccine supply			
Wrong vaccine	Records	Out of date		1
	inaccurate/	records		
	unclear			
Wrong vaccine	Out of date			1
	records			
Wrong vaccine	Vaccine			1
	prescribing and			
	dispensing			
	incident			
Wrong vaccine	Wrong vaccine	Records		1
J S CALLSTON	3	inaccurate/		
		accarace/		

		unclear		
Wrong vaccine	Wrong timing			1
Wrong dose	Non-specific	Appointment		1
	vaccination-	management		
	related incident			
Wrong dose	Reconstitution			1
	error			
Wrong dose	Wrong patient			1
Wrong timing	Communication -	Appointment		1
	professionals	management		
Wrong timing	Communication -	Appointment		1
	patients/	management		
	caregivers			
Wrong timing	Communication -	Out of date		1
	patients/	records		
	caregivers			
Wrong timing	Insufficient			1
	vaccine supply			
Wrong timing	Other	Appointment		1
	administrative	management		
	incidents			
Wrong timing	Reconstitution			1
	error			
Wrong timing	Record	Appointment		1
	availability	management		
Wrong timing	Records	Out of date	Communication -	1
	inaccurate/	records	patients/	
	unclear		caregivers	
Wrong timing	Out of date	Communication -		1
	records	professionals		
Wrong timing	Out of date	Transfer of		1
	records	information		
Wrong timing	Transfer of	Communication -		1
	information	patients/		
		caregivers		
Wrong timing	Transfer of	Out of date		1
	information	records		
Wrong timing	Transfer of			1
	information			

Wrong timing	Vaccine not			1
	administered			
Wrong timing	Vaccine not	Appointment		1
	administered	management		
Wrong timing	Vaccine not	Record	Insufficient	1
	administered	availability	vaccine supply	
Wrong timing	Wrong vaccine	Communication -		1
		patients/		
		caregivers		
Wrong timing	Wrong vaccine	Records		1
		inaccurate/		
		unclear		
Wrong timing	Wrong vaccine	Out of date		1
		records		
				1745

3.2 The frequencies of combinations of contributory factors for the primary incident types related to vaccination

	Con	tributory factors	Frequency of
Primary incident type			combination
Wrong number of doses			269
Wrong timing			186
Wrong vaccine			123
Adverse reaction			122
Records not up to date			51
Procedural error	Non-compliance		39
	Non-specific		
Wrong number of doses	mistake		36
	Failure to follow		
Wrong number of doses	protocol		35
Not administered			33
Communication with			
patient/ caregiver			31
Appointment			
management			26
Inaccurate records			26
Out of date vaccine			26
	Patient/ caregiver		
Wrong number of doses	knowledge		26
	Similar vaccine		
Wrong vaccine	names		25
	Similar vaccine		
Wrong number of doses	names		23
	Non-specific		
Wrong vaccine	mistake		21
Prescribing/dispensing			
error			19
Adverse reaction	Patient allergy		18
	Failure to follow		
Wrong vaccine	protocol		18
Wrong dose			16
	Failure to follow		
Wrong timing	protocol		16

	Vaccine/		
Wrong dose	equipment failure		14
	Non-specific		
Wrong timing	mistake		13
Wrong dose	Non-compliance		11
Non-specific			
administration error			10
	Vaccine/		
Not administered	equipment design		10
	Vaccine/		
Procedural error	equipment failure		10
Wrong number of doses	Continuity of care		8
	Failure to follow		
Out of date vaccine	protocol		7
Procedural error			7
Transfer of records	Continuity of care		7
Transfer of records			7
Wrong number of doses	Misreading		7
Wrong timing	Continuity of care		7
Wrong timing	Working conditions		7
Records not available			6
Records not up to date	New to area		6
Wrong timing	Continuity of care		6
Communication with			
patient/ caregiver			5
Environmental hazard			5
Non-specific	Failure to follow		
administration error	protocol		5
Wrong number of doses	Non-disclosure		5
Wrong timing	Continuity of care	New to area	5
Communication with			
patient/ caregiver			4
Communication with	Non-specific		
patient/ caregiver	mistake		4
Contraindicated vaccine			4
	Non-specific	Vaccine/	
	mistake	equipment	
Not administered		packaging	4
Not administered	Vaccine/		4

	equipment failure		
Not administered	Staff distraction		4
	Vaccine/		
Out of date vaccine	equipment storage		4
Vaccine recall			4
	Failure to follow		
Wrong dose	protocol		4
	Similar patient		
Wrong patient	names		4
	Failure to follow	Education and	
Wrong timing	protocol	training	4
Wrong timing	New to area		4
Wrong vaccine	Staff distraction		4
Appointment	Non-specific		
management	mistake		3
Communication with	Failure to follow		
patient/ caregiver	protocol		3
	Non-specific		
Not administered	mistake		3
	Vaccine/		
Not administered	equipment design		3
Reconstitution error			3
Referral			3
	Non-specific		
Wrong dose	mistake		3
	Patient/ caregiver	Failure to follow	
Wrong number of doses	knowledge	protocol	3
	Non-specific	Failure to follow	
Wrong number of doses	mistake	protocol	3
Wrong number of doses	Looked-after child		3
	New to area	Looked-after	
Wrong number of doses		child	3
	New to area	Continuity of	
Wrong number of doses		care	3
Wrong patient			3
	Failure to follow		
Wrong site	protocol		3
	Failure to follow		
Wrong timing	protocol		3

Wrong vaccine	Working conditions		3
	Similar vaccine	Vaccine/	
	names	equipment	
Wrong vaccine		storage	3
	Non-specific	Failure to follow	
Wrong vaccine	mistake	protocol	3
	Non-specific	Inadequate	
	mistake	guidelines or	
Wrong vaccine		protocols	3
	Non-specific	Working	
Wrong vaccine	mistake	conditions	3
	Non-specific	Vaccine/	
	mistake	equipment	
Wrong vaccine		storage	3
	Non-specific	Non-disclosure	
Adverse reaction	patient health		2
Adverse reaction	Non-compliance		2
Appointment	New to area		
management			2
Appointment	Working conditions		
management			2
Appointment	Working conditions		
management			2
Appointment	Failure to follow		
management	protocol		2
Communication between			
professionals			2
	Non-specific		
Contraindicated vaccine	mistake		2
Contraindicated vaccine	Non-disclosure		2
Contraindicated vaccine	Patient allergy		2
	Failure to follow		
Inaccurate records	protocol		2
Non-specific	Non-specific		
administration error	mistake		2
Not administered	Non-compliance		2
Not administered	Staff knowledge		2
	Failure to follow	Other staff	
Not administered	protocol	factors	2

	Staff distraction	Working		
Not administered		conditions		2
	Similar vaccine	Vaccine/	Vaccine/	
	names	equipment	equipment	
Not administered		packaging	storage	2
	Failure to follow	Inadequate		
	protocol	guidelines or		
Out of date vaccine		protocols		2
Out of date vaccine	Misreading			2
	Non-specific	Working		
Out of date vaccine	mistake	conditions		2
	Similar vaccine	Vaccine/	Vaccine/	
Prescribing/dispensing	names	equipment	equipment	
error		storage	packaging	2
Records not up to date	Working conditions			2
Used needle				2
Wrong dose	Staff distraction			2
	Patient/ caregiver	New to area		
Wrong number of doses	knowledge			2
	Failure to follow	Non-specific		
Wrong number of doses	protocol	mistake		2
Wrong number of doses	Inattention			2
Wrong number of doses	Staff distraction			2
	Similar patient			
Wrong number of doses	names			2
	Non-specific	Working		
Wrong number of doses	mistake	conditions		2
Wrong number of doses	Other staff factors			2
	Inadequate			
	guidelines or			
Wrong number of doses	protocols			2
Wrong number of doses	Continuity of care			2
Wrong number of doses	Working conditions			2
	New to area	Patient/		
		caregiver		
Wrong number of doses		knowledge		2
	Failure to follow	Education and		
Wrong number of doses	protocol	training		2
Wrong number of doses	Failure to follow	Working		2

	protocol	conditions		
	Non-specific			
Wrong patient	mistake			2
Wrong site				2
	Vaccine/			
Wrong storage	equipment failure			2
	Staff knowledge	Education and		
Wrong timing		training		2
Wrong timing	Patient language			2
	Non-specific	Failure to follow		
Wrong timing	mistake	protocol		2
	Patient/ caregiver			
Wrong timing	knowledge			2
	Failure to follow	Working		
Wrong timing	protocol	conditions		2
	Inadequate			
	guidelines or			
Wrong timing	protocols			2
	Continuity of care	Looked-after		
Wrong timing		child		2
	Vaccine/			
Wrong timing	equipment failure			2
	Similar vaccine	Vaccine/	Vaccine/	
	names	equipment	equipment	
Wrong vaccine		packaging	storage	2
Wrong vaccine	Misreading			2
Wrong vaccine	Staff knowledge			2
	Education and			
Wrong vaccine	training			2
	Failure to follow	Education and		
Wrong vaccine	protocol	training		2
	Inadequate	Continuity of		
	guidelines or	care		
Access to care	protocols			1
	Abnormal			
Adverse reaction	coagulation			1
	Failure to follow			
Adverse reaction	protocol			1
Appointment	Continuity of care			1

management				
Appointment	Looked-after child			
management				1
Communication with	Failure to follow	Staff distraction		
patient/ caregiver	protocol			1
Communication with	Patient/ caregiver			
patient/ caregiver	knowledge			1
Communication with	Working conditions			
patient/ caregiver				1
	Inadequate	Looked-after		
Communication with	guidelines or	child		
patient/ caregiver	protocols			1
	Staff knowledge	Working		
Contraindicated vaccine		conditions		1
	Immunocompromis	Failure to follow		
Contraindicated vaccine	ed patient	protocol		1
	Non-specific	Immunocompro		
Contraindicated vaccine	mistake	mised patient		1
	Pregnancy	Patient/		
	3,	caregiver		
Contraindicated vaccine		knowledge		1
	Non-specific	5		
Contraindicated vaccine	patient health			1
	Immunocompromis			
Contraindicated vaccine	ed patient			1
Contramareacea vaccine	Abnormal			•
Contraindicated vaccine	coagulation			1
Contraindicated vaccine	Staff knowledge			1
contramareacea vaccine	Failure to follow			
Contraindicated vaccine	protocol			1
Contraindicated vaccine	Inattention			1
Inaccurate records	Working conditions			1
	Looked-after child			
Inaccurate records				1
1	Non-specific			4
Inaccurate records	mistake			1
Insufficient supply	5		11.	1
	Patient allergy	Failure to follow	Working	
Medication error		protocol	conditions	1
Non-specific	Inattention			1

administration error				
Non-specific	Vaccine/			
administration error	equipment storage			1
Non-specific	Non-disclosure			
administration error				1
Non-specific	Failure to follow	Looked-after		
administration error	protocol	child		1
Non-specific	New to area			
administration error				1
Non-specific	Patient/ caregiver			
administration error	knowledge			1
Non-specific	Vaccine/			
administration error	equipment failure			1
Non-specific	Working conditions			
administrative error				1
Non-specific vaccine	Vaccine/			
error	equipment storage			1
	Other staff factors	Working		
Not administered		conditions		1
Not administered	Working conditions			1
	Vaccine/	Failure to follow		
	equipment	protocol		
Not administered	packaging			1
	Inadequate	Vaccine/		
	guidelines or	equipment		
Not administered	protocols	packaging		1
	Inadequate			
	guidelines or			
Not administered	protocols			1
Not administered	Working conditions			1
Not administered	Patient disability	Violence		1
	Failure to follow			
Not administered	protocol			1
	Non-specific	Working		
Not administered	mistake	conditions		1
Out of date vaccine	Inattention			1
	Non-specific	Failure to follow	Vaccine/	
	mistake	protocol	equipment	
Out of date vaccine			storage	1

	Failure to follow	Non-specific		
Out of date vaccine	protocol	mistake		1
Out of date vaccine				1
	Non-specific	Vaccine/	Vaccine/	
	mistake	equipment	equipment	
Out of date vaccine		packaging	storage	1
	Non-specific	Working		
Out of date vaccine	mistake	conditions		1
Prescribing / dispensing	Non-specific			
error	mistake			1
	Similar vaccine	Failure to follow	Vaccine/	
Prescribing / dispensing	names	protocol	equipment	
error			storage	1
	Similar vaccine	Vaccine/	Working	
Prescribing / dispensing	names	equipment	conditions	
error		storage		1
Prescribing / dispensing	Vaccine/			
error	equipment storage			1
Prescribing/ dispensing	Non-specific			
incident	mistake			1
	Staff knowledge	Working		
Procedural error		conditions		1
Professionalism				1
	Other staff factors	Inadequate		
		guidelines or		
Professionalism		protocols		1
	Failure to follow	Staff knowledge		
Reconstitution error	protocol			1
	Inadequate			
	guidelines or			
Records not up to date	protocols			1
	New to area	Looked-after		
Records not up to date		child		1
	Patient/ caregiver			
Records not up to date	knowledge			1
	Continuity of care	New to area	Looked-after	
Records not up to date			child	1
	Failure to follow			
Records not up to date	protocol			1

	Inadequate			
	guidelines or			
Records not up to date	protocols			1
Records not up to date	Continuity of care			1
Transfer of records	New to area			1
Transport error	Patient allergy			1
Used needle	Non-compliance			1
	Failure to follow	Non-compliance		
Used needle	protocol			1
	Non-specific			
Used needle	mistake			1
Wrong dose	Staff knowledge			1
	Vaccine/	Non-compliance		
Wrong dose	equipment failure			1
	Inadequate			
	guidelines or			
Wrong dose	protocols			1
	Education and	Inadequate		
	training	guidelines or		
Wrong dose		protocols		1
	Misreading	Working		
Wrong dose		conditions		1
	Non-specific	Failure to follow		
Wrong dose	mistake	protocol		1
	Similar vaccine			
Wrong dose	names			1
	Misreading	Looked-after		
Wrong number of doses		child		1
	Vaccine/	Failure to follow		
	equipment	protocol		
Wrong number of doses	packaging			1
	Non-specific	Failure to follow	Working	
Wrong number of doses	mistake	protocol	conditions	1
	Similar patient	Working		
Wrong number of doses	names	conditions		1
	Inadequate	Continuity of		
	guidelines or	care		
Wrong number of doses	protocols			1
Wrong number of doses	Working conditions			1

	Similar patient	Failure to follow		
Wrong number of doses	names	protocol		1
	Non-specific	Patient/		
	mistake	caregiver		
Wrong number of doses		knowledge		1
	Non-specific	Failure to follow	Patient/	
	mistake	protocol	caregiver	
Wrong number of doses			knowledge	1
	Non-specific	Failure to follow	Education	
Wrong number of doses	mistake	protocol	and training	1
	Education and			
Wrong number of doses	training			1
	New to area	Patient	Failure to	
		language	follow	
Wrong number of doses			protocol	1
	New to area	Patient/	Continuity of	
		caregiver	care	
Wrong number of doses		knowledge		1
Wrong number of doses	Non-compliance			1
	Patient/ caregiver	Misreading		
Wrong number of doses	knowledge			1
Wrong number of doses	Staff knowledge			1
	Staff knowledge	Education and		
Wrong number of doses		training		1
	Failure to follow			
Wrong number of doses	protocol			1
	Failure to follow	Misreading		
Wrong number of doses	protocol			1
	Failure to follow	Patient/		
	protocol	caregiver		
Wrong number of doses		knowledge		1
	Failure to follow	Patient/	New to area	
	protocol	caregiver		
Wrong number of doses		knowledge		1
	Failure to follow	Looked-after		
Wrong number of doses	protocol	child		1
	Failure to follow	Similar vaccine		
Wrong number of doses	protocol	names		1
Wrong number of doses	Failure to follow	Other staff	Staff	1

	protocol	factors	knowledge	
	Inattention	Failure to follow		
Wrong number of doses		protocol		1
	Patient/ caregiver	Working		
Wrong number of doses	knowledge	conditions		1
Wrong patient	Non-disclosure			1
	Failure to follow			
Wrong patient	protocol			1
Wrong patient	Staff distraction			1
	Similar patient	Patient		
Wrong patient	names	disability		1
	Non-specific	Patient		
Wrong patient	mistake	disability		1
	Inadequate			
	guidelines or			
Wrong patient	protocols			1
Wrong patient	Working conditions			1
Wrong route				1
	Failure to follow			
Wrong route	protocol			1
Wrong route	Misreading			1
	Misreading	Working	Vaccine/	
		conditions	equipment	
Wrong site			failure	1
Wrong site	Non-compliance			1
	Failure to follow	Vaccine/		
	protocol	equipment		
Wrong storage		failure		1
Wrong timing	Access difficulties			1
	Failure to follow	Other staff		
Wrong timing	protocol	factors		1
Wrong timing	Working conditions			1
Wrong timing	Patient disability			1
	Caregiver/ patient	Access		
Wrong timing	behaviour	difficulties		1
	Other staff factors	Staff knowledge	Education	
Wrong timing			and training	1
	Inadequate			
Wrong timing	guidelines or			1

	protocols			
	Continuity of care	Patient/		
		caregiver		
Wrong timing		knowledge		1
Wrong timing	New to area			1
	Patient language	Failure to follow		
Wrong timing		protocol		1
Wrong timing	Non-disclosure			1
	Failure to follow	New to area		
Wrong timing	protocol			1
	Failure to follow	Non-specific		
Wrong timing	protocol	mistake		1
	Failure to follow	Staff distraction		
Wrong timing	protocol			1
	Failure to follow	Education and		
Wrong timing	protocol	training		1
<u> </u>	Staff distraction	Working		
Wrong timing		conditions		1
Wrong timing	Misreading			1
3 3	Non-specific	Patient		
Wrong timing	mistake	language		1
3 3	Non-specific	Patient/		
	mistake	caregiver		
Wrong timing		knowledge		1
3 3	Non-specific	Education and	New to area	
Wrong timing	mistake	training		1
Wrong timing	Working conditions	3		1
	Education and			
Wrong timing	training			1
	Education and	Other staff		
Wrong timing	training	factors		1
Wrong timing	Non-compliance			1
	Non-specific			•
Wrong timing	patient health			1
Wrong timing	Looked-after child			1
Wrong timing	Continuity of care	New to area		1
THOUS CHIMIS	Failure to follow	Staff knowledge	Education	1
Wrong vaccine	protocol	Juli Kilowieuge	and training	1
Wrong vaccine	Staff knowledge	Education and	and training	1
WI OIIS VACCINE	Stall knowledge	Luucation and		I

		training		
	Other staff factors	Failure to follow		
Wrong vaccine		protocol		1
	Staff fatigue/			
Wrong vaccine	stress			1
	Staff fatigue/	Working		
Wrong vaccine	stress	conditions		1
	Failure to follow	Other staff		
Wrong vaccine	protocol	factors		1
	Vaccine/	Non-specific	Failure to	
	equipment failure	mistake	follow	
Wrong vaccine			protocol	1
	Vaccine/			
	equipment			
Wrong vaccine	packaging			1
Wrong vaccine	Working conditions			1
	Working conditions	Failure to follow	Looked-after	
Wrong vaccine		protocol	child	1
	Working conditions	Education and		
Wrong vaccine		training		1
	Education and	New to area	Similar	
	training		vaccine	
Wrong vaccine			names	1
	Vaccine/			
Wrong vaccine	equipment storage			1
	Failure to follow	Staff knowledge		
Wrong vaccine	protocol			1
Wrong vaccine	Inattention	Staff distraction		1
	Similar vaccine	Failure to follow	Vaccine/	
	names	protocol	equipment	
Wrong vaccine			storage	1
	Similar vaccine	Staff fatigue/		
Wrong vaccine	names	stress		1
Wrong vaccine	Pregnancy			1
Wrong vaccine	Staff knowledge			1
	Failure to follow	Non-specific		
Wrong vaccine	protocol	mistake		1
	Failure to follow	Inattention		
Wrong vaccine	protocol			1

	Failure to follow	Working	Education	
Wrong vaccine	protocol	conditions	and training	1
Wrong vaccine	Inattention			1
	Inattention	Similar vaccine		
Wrong vaccine		names		1
	Staff distraction	Staff fatigue/	Staff	
		stress	fatigue/	
Wrong vaccine			stress	1
	Staff distraction	Working		
Wrong vaccine		conditions		1
	Staff distraction	Working	Education	
Wrong vaccine		conditions	and training	1
	Similar vaccine	Failure to follow		
Wrong vaccine	names	protocol		1
	Similar vaccine	Similar vaccine		
Wrong vaccine	names	names		1
Total				1745

3.3 The frequency of each combination of incidents involving 'unwell' children

Drimany incident		Frequency of	
Primary incident		Contributory incidents	combination
Medication dispensing			283
Transfer of information			88
Communication -			
patients / caregivers			87
Inadequate triaging			84
Treatment and			
procedures			82
Equipment			66
Medication prescribing			54
	Inadequate		
Inadequate triaging	history		48
Documentation			45
	Medication		
Medication dispensing	prescribing		44
Medication	Medication		
administration	dispensing		35
Other			33
Investigations			32
Medication			
administration			32
Access to care			23
Incomplete referral			23
Transfer of patients			23
	Medication		
Medication dispensing	dispensing		22
Treatment decision			22
	Communication		
	- patients /		
Inadequate triaging	caregivers		21
	Inadequate		
Delayed referral	triaging		19
Communication			
between professionals			18

procedures December December	Treatment and	Equipment		
Delayed assessment Diagnosis Inadequate history Other medication Delayed referral Delayed r	procedures			17
Diagnosis 14 14 14 14 14 14 14 1	Inadequate triaging	Documentation		16
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Inadequate discharge 8	Delayed assessment	Access to care		8
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planning				
Referral administrative				
issues				8
Communication -	Inadequate			
patients / caregivers	triaging			7
	Failure to			
	identify at risk/			
Delayed referral	vulnerable child			7
Failure to refer when	Inadequate			
appropriate	triaging			7
Insufficient assessment				,
(non-specific)				7
Medication	Medication			,
administration	prescribing			7
Other medication	Medication			,
Other medication				7
Communication	dispensing			,
Communication -	Inadequate			
patients / caregivers	history			6
	Inadequate			
Documentation	history			6
Equipment	Equipment			6
	Other			
Equipment	administrative			6
Failure to identify at				
risk/ vulnerable child				6
	Inadequate	Documentation		
Inadequate triaging	history			6
	Failure to			
	identify at risk/			
Inadequate triaging	vulnerable child			6
	Transfer of			
Transfer of information	information			6
Treatment and	Equipment			
procedures				6
	Appointment			
Access to care	management			5
Communication -	Transfer of			
patients / caregivers	information			5
Communication -	Failure to			5

patients / caregivers	identify at risk/			
	vulnerable child			
	Transfer of			
Delayed assessment	information			5
	Inadequate	Inadequate	Communi	
	triaging	history	cation -	
			patients /	
Delayed referral			caregivers	5
	Inadequate	Communication -		
	triaging	patients /		
Delayed referral		caregivers		5
	Delayed			
Delayed referral	assessment			5
	Communication			
	- patients /			
Documentation	caregivers			5
	Documentation	Inadequate		
Inadequate triaging		history		5
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Incomplete referral	triaging			5
Medication	Medication	Medication		
administration	dispensing	prescribing		5
Medication	Medication	Medication		
administration	dispensing	dispensing		5
Medication	Equipment			
administration				5
	Communication			
	- patients /			
Medication dispensing	caregivers			5
Other diagnosis and				
assessment				5
	Communication			
	between			
Transfer of information	professionals			5
Transfer of information	Equipment			5
Communication -	Investigations			
patients / caregivers				4
	Failure to			
Delayed referral	identify at risk/			4

Inadequate discharge planning vulnerable child vulnerable		vulnerable child			
planning vulnerable child Communication - patients / caregivers		Failure to			
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Inadequate examination 3	appropriate				3
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	Documentation	Inadequate	Communi	
		history	cation -	
			patients /	
Inadequate triaging			caregivers	3
Insufficient assessment	Equipment			
(non-specific)				3
	Communication			
	- patients /			
Investigations	caregivers			3
	Medication	Medication		
Medication dispensing	dispensing	prescribing		3
Other administrative				3
	Communication			
	between			
Transfer of information	professionals			3
	Appointment			
Transfer of information	management			3
Transfer of patients	Equipment			3
	Insufficient			
	assessment			
Transfer of patients	(non-specific)			3
Treatment and	Treatment and			
procedures	procedures			3
	Medication			
Treatment decision	dispensing			3
Treatment decision	Diagnosis			3
	Incomplete			
Access to care	referral			2
	Other			
Access to care	administrative			2
	Incomplete			
Access to care	referral			2
	Communication			
Appointment	between			
management	professionals			2
	Failure to refer			
Communication -	when			
patients / caregivers	appropriate			2
Communication -	Incomplete			2

patients / caregivers	referral			
	Diagnosis	Insufficient		
Communication -		assessment (non-		
patients / caregivers		specific		2
Communication -	Inadequate	Inadequate		
patients / caregivers	triaging	history		2
	Inadequate	Communication		
Communication -	triaging	between		
patients / caregivers		professionals		2
	Documentation	Transfer of		
Delayed assessment		information		2
	Referral			
	administrative			
Delayed assessment	issues			2
	Inadequate	Communication -		
	triaging	patients /		
Delayed assessment		caregivers		2
	Inadequate	Appointment	Communi	
	discharge	management	cation -	
	planning		patients /	
Delayed assessment			caregivers	2
	Transfer of			
Delayed referral	information			2
	Inadequate	Failure to identify		
	triaging	at risk/		
Delayed referral		vulnerable child		2
	Communication			
	between			
Delayed referral	professionals			2
	Transfer of			
Delayed referral	patients			2
Delayed referral	Documentation			2
	Communication			
	- patients /			
Delayed referral	caregivers			2
Delayed referral	Equipment			2
	Investigations	Failure to identify		
		at risk/		
Diagnosis		vulnerable child		2

	Other			
Documentation	administrative			2
Documentation	Investigations			2
	Inadequate			
	discharge			
Equipment	planning			2
Failure to refer when	Inadequate			
appropriate	history			2
Failure to refer when	Transfer of			
appropriate	information			2
	Failure to			
	identify at risk/			
Inadequate history	vulnerable child			2
	Communication			
	- patients /			
Inadequate triaging	caregivers			2
	Inadequate	Documentation	Communi	
	history		cation -	
			patients /	
Inadequate triaging			caregivers	2
	Failure to	Communication -		
	identify at risk/	patients /		
Inadequate triaging	vulnerable child	caregivers		2
	Failure to			
	identify at risk/			
Incomplete referral	vulnerable child			2
Insufficient assessment	Treatment and			
(non-specific)	procedures			2
	Transfer of	Appointment		
Investigations	information	management		2
Medication	Transfer of			
administration	information			2
	Communication			
	between			
Medication dispensing	professionals			2
	Communication			
	between			
Medication dispensing	professionals			2
Medication monitoring	Appointment			2

	management			
Medication prescribing	Documentation			2
	Appointment			
Other	management			2
Other	Other			2
	Transfer of			
Other	information			2
Other	Access to care			2
Other medication	Equipment			2
	Communication			
	- patients /			
Treatment decision	caregivers			2
	Transfer of			
Treatment decision	information			2
	Inadequate			
Treatment decision	examination			2
	Medication			
Treatment decision	prescribing			2
	Other	Communication -		
	administrative	patients /		
Access to care		caregivers		1
	Transfer of	Equipment		
Access to care	information			1
	Transfer of	Transfer of		
Access to care	information	information		1
	Transfer of	Other		
Access to care	information			1
	Inadequate	Inadequate		
Access to care	triaging	history		1
	Communication	Documentation		
	- patients /			
Access to care	caregivers			1
	Appointment	Inadequate	Failure to	
	management	triaging	identify	
			at risk/	
			vulnerabl	
Access to care			e child	1
Access to care	Equipment			1
Access to care	Transfer of	Transfer of		1

	information	information		
	Appointment	Inadequate		
Access to care	management	triaging		1
	Transfer of			
Access to care	patients			1
	Appointment	Failure to refer		
Access to care	management	when appropriate		1
	Transfer of	Communication		
	patients	between		
Access to care		professionals		1
	Referral			
	administrative			
Access to care	issues			1
	Inadequate			
Access to care	triaging			1
	Referral			
Appointment	administrative			
management	issues			1
	Failure to	Inadequate		
Appointment	identify at risk/	triaging		
management	vulnerable child			1
	Transfer of	Inadequate		
Communication	information	discharge		
between professionals		planning		1
Communication	Documentation			
between professionals				1
Communication	Inadequate			
between professionals	history			1
Communication -	Treatment			
patients / caregivers	decision			1
Communication -	Medication			
patients / caregivers	prescribing			1
	Inadequate	Access to care	Appointm	
	triaging		ent	
Communication -			managem	
patients / caregivers			ent	1
Communication -	Other			
patients / caregivers				1
Communication -	Other	Documentation		1

patients / caregivers				
Communication -	Documentation	Inadequate		
patients / caregivers		history		1
Communication -	Delayed referral	Inadequate		
patients / caregivers		triaging		1
	Failure to refer	Failure to identify		
Communication -	when	at risk/		
patients / caregivers	appropriate	vulnerable child		1
Communication -	Incomplete	Inadequate		
patients / caregivers	referral	triaging		1
	Diagnosis	Failure to identify		
Communication -		at risk/		
patients / caregivers		vulnerable child		1
Communication -	Inadequate	Inadequate	Other	
patients / caregivers	triaging	history		1
	Failure to	Equipment		
Communication -	identify at risk/			
patients / caregivers	vulnerable child			1
	Failure to	Inadequate		
Communication -	identify at risk/	triaging		
patients / caregivers	vulnerable child			1
	Insufficient	Access to care		
Communication -	assessment			
patients / caregivers	(non-specific			1
	Insufficient	Equipment	Equipmen	
Communication -	assessment		t	
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Communication -	Delayed	Inadequate		
patients / caregivers	assessment	history		1
Communication -	Treatment and			
patients / caregivers	procedures			1
Communication -	Treatment	Inadequate		
patients / caregivers	decision	examination		1
Communication -	Equipment			
patients / caregivers				1
Communication -	Appointment			
patients / caregivers	management			1
Communication -	Inadequate			
Communication	madequate			

	Other			
Delayed assessment	administrative			1
	Appointment	Transfer of		
Delayed assessment	management	information		1
	Access to care	Appointment		
Delayed assessment		management		1
	Access to care	Referral		
		administrative		
Delayed assessment		issues		1
Delayed assessment	Access to care	Equipment		1
	Access to care	Inadequate		
Delayed assessment		triaging		1
	Other	Appointment		
Delayed assessment		management		1
Delayed assessment	Documentation			1
Delayed assessment	Delayed referral			1
	Delayed referral	Failure to identify		
		at risk/		
Delayed assessment		vulnerable child		1
	Failure to refer			
	when			
Delayed assessment	appropriate			1
	Referral	Communication		
	administrative	between		
Delayed assessment	issues	professionals		1
	Inadequate	Documentation	Document	
Delayed assessment	triaging		ation	1
	Inadequate	Inadequate		
Delayed assessment	triaging	history		1
	Inadequate	Inadequate	Communi	
	triaging	history	cation -	
			patients /	
Delayed assessment			caregivers	1
	Inadequate	Failure to identify		
	triaging	at risk/		
Delayed assessment		vulnerable child		1
	Inadequate	Communication		
	triaging	between		
Delayed assessment		professionals		1

	Inadequate	Equipment			
Delayed assessment	triaging				1
	Failure to				
	identify at risk/				
Delayed assessment	vulnerable child				1
	Failure to	Communication			
	identify at risk/	between			
Delayed assessment	vulnerable child	professionals			1
Delayed assessment	Investigations	Access to care			1
	Communication	Inadequate			
	- patients /	triaging			
Delayed assessment	caregivers				1
	Communication	Inadequate	Inadequat		
	- patients /	triaging	e triaging		
Delayed assessment	caregivers				1
	Communication	Incomplete	Access to	Appoin	
	between	referral	care	tment	
	professionals			manag	
Delayed assessment				ement	1
	Communication	Incomplete	Failure to		
	between	referral	identify		
	professionals		at risk/		
			vulnerabl		
Delayed assessment			e child		1
Delayed assessment	Equipment				1
	Failure to	Transfer of			
	arrange follow	information			
Delayed referral	up				1
	Incomplete	Failure to identify			
	referral	at risk/			
Delayed referral		vulnerable child			1
Delayed referral	Diagnosis	Documentation			1
Delayed referral	Diagnosis	Diagnosis			1
Delayed referral	Access to care				1
Delayed referral	Other				1
	Diagnosis	Incomplete			
Delayed referral		referral			1
	Insufficient				
Delayed referral	assessment				1

	(non-specific)				
	Inadequate	Access to care			
Delayed referral	triaging				1
	Inadequate	Documentation	Inadequat		
Delayed referral	triaging		e history		1
	Inadequate	Documentation	Inadequat	Comm	
	triaging		e history	unicati	
				on -	
				patien	
				ts /	
				caregi	
Delayed referral				vers	1
	Inadequate	Medication			
Delayed referral	triaging	dispensing			1
	Inadequate				
Delayed referral	history				1
	Inadequate	Failure to identify	Communi		
	examination	at risk/	cation -		
		vulnerable child	patients /		
Delayed referral			caregivers		1
	Delayed	Access to care			
Delayed referral	assessment				1
	Delayed	Inadequate			
Delayed referral	assessment	triaging			1
	Delayed	Inadequate	Document		
Delayed referral	assessment	triaging	ation		1
	Delayed	Inadequate	Communi		
	assessment	triaging	cation		
			between		
			profession		
Delayed referral			als		1
	Treatment	Communication -	Failure to	Inadeq	
	decision	patients /	identify	uate	
		caregivers	at risk/	exami	
			vulnerabl	nation	
Delayed referral			e child		1
	Investigations	Failure to identify			
		at risk/			
Delayed referral		vulnerable child			1

	Communication	Inadequate		
	- patients /	triaging		
Delayed referral	caregivers			1
	Communication	Inadequate	Failure to	
	- patients /	triaging	identify	
	caregivers		at risk/	
			vulnerabl	
Delayed referral			e child	1
	Inadequate			
Delayed referral	triaging			1
	Communication			
	- patients /			
Delayed referral	caregivers			1
Delayed referral	Diagnosis	Documentation		1
	Failure to			
	identify at risk/			
Diagnosis	vulnerable child			1
	Investigations	Communication		
		between		
Diagnosis		professionals		1
	Transfer of			
Diagnosis	information			1
	Failure to	Inadequate	Transfer	
	identify at risk/	history	of	
	vulnerable child		informati	
Diagnosis			on	1
	Failure to	Investigations		
	identify at risk/			
Diagnosis	vulnerable child			1
Diagnosis	Delayed referral	Documentation		1
	Investigations	Diagnosis	Investigat	
Diagnosis			ions	1
Diagnosis	Access to care			1
	Delayed referral	Failure to identify		
		at risk/		
Diagnosis		vulnerable child		1
Diagnosis	Diagnosis			1
	Appointment	Communication -		
Diagnosis	management	patients /		1

		caregivers		
	Transfer of	Other	Other	
	information	administrative	administr	
Diagnosis			ative	1
Diagnosis	Delayed referral			1
	Inadequate			
Diagnosis	examination			1
	Failure to			
	identify at risk/			
Diagnosis	vulnerable child			1
	Investigations	Inadequate		
Diagnosis		history		1
	Investigations	Communication		
		between		
Diagnosis		professionals		1
	Investigations	Other		
Diagnosis		administrative		1
	Communication			
	- patients /			
Diagnosis	caregivers			1
	Transfer of			
Documentation	information			1
	Appointment			
Documentation	management			1
Documentation	Equipment			1
	Communication	Transfer of		
	between	information		
Documentation	professionals			1
Documentation	Investigations			1
Documentation	Other	Other		1
	Insufficient			
1	assessment			
Documentation	(non-specific)			1
	Inadequate	Communication -		
	history	patients /		
Documentation		caregivers		1
	Inadequate	Communication -	Communi	
	history	patients /	cation -	
Documentation		caregivers	patients /	1

			caregivers		
	Inadequate				
Documentation	examination				1
	Insufficient	Appointment			
	assessment	management			
Documentation	(non-specific)				1
	Communication	Inadequate			
	- patients /	history			
Documentation	caregivers				1
	Medication				
Equipment	dispensing				1
	Equipment	Referral	Incomplet		
		administrative	e referral		
Equipment		issues			1
	Access to care	Transfer of			
Equipment		information			1
	Failure to refer				
	when				
Equipment	appropriate				1
	Inadequate	Transfer of			
	discharge	information			
Equipment	planning				1
	Medication				
Equipment	prescribing				1
	Other				
Equipment	medication				1
	Transfer of				
Equipment	information				1
	Delayed				
Equipment	assessment				1
Failure to arrange	Delayed				
follow up	assessment				1
Failure to identify at	Inadequate	Inadequate			
risk/ vulnerable child	triaging	history			1
Failure to refer when	Investigations	Investigations			
appropriate					1
Failure to refer when	Access to care	Appointment			
appropriate		management			1
Failure to refer when	Transfer of	Communication			1

appropriate	information	between		
		professionals		
	Inadequate	Inadequate	Communi	
	triaging	history	cation -	
Failure to refer when			patients /	
appropriate			caregivers	1
	Inadequate	Failure to identify		
Failure to refer when	triaging	at risk/		
appropriate		vulnerable child		1
	Communication			
Failure to refer when	- patients /			
appropriate	caregivers			1
	Communication	Failure to identify		
Failure to refer when	- patients /	at risk/		
appropriate	caregivers	vulnerable child		1
	Communication			
Failure to refer when	between			
appropriate	professionals			1
Inadequate discharge	Transfer of	Access to care		
planning	information			1
Inadequate discharge	Documentation			
planning				1
	Failure to refer			
Inadequate discharge	when			
planning	appropriate			1
Inadequate discharge	Incomplete			
planning	referral			1
	Communication	Transfer of		
Inadequate discharge	between	information		
planning	professionals			1
	Communication	Transfer of		
Inadequate discharge	between	information		
planning	professionals			1
Inadequate discharge	Equipment			
planning				1
	Equipment	Medication	Communi	
		dispensing	cation -	
Inadequate discharge			patients /	
planning			caregivers	1

	Transfer of	Transfer of			
Inadequate examination	information	information			1
	Documentation	Transfer of			
Inadequate examination		information			1
Inadequate examination	Equipment				1
	Other				
Inadequate triaging	administrative				1
	Transfer of				
Inadequate triaging	information				1
	Documentation	Inadequate	Communi	Other	
		history	cation -		
			patients /		
Inadequate triaging			caregivers		1
	Inadequate				
Inadequate triaging	triaging				1
	Inadequate	Transfer of			
Inadequate triaging	history	information			1
	Inadequate	Failure to identify			
	history	at risk/			
Inadequate triaging		vulnerable child			1
	Inadequate	Communication -	Other		
	history	patients /			
Inadequate triaging		caregivers			1
	Inadequate	Communication -	Document		
	history	patients /	ation		
Inadequate triaging		caregivers			1
	Inadequate	Communication			
	history	between			
Inadequate triaging		professionals			1
	Medication				
Inadequate triaging	dispensing				1
	Communication	Documentation			
	between				
Inadequate triaging	professionals				1
Inadequate triaging	Equipment				1
	Appointment				
Incomplete referral	management				1
	Insufficient				
Incomplete referral	assessment				1

	(non-specific)				
	Inadequate				
Incomplete referral	examination				1
	Communication	Inadequate	Delayed	Other	
	- patients /	history	assessme		
Incomplete referral	caregivers		nt		1
	Incomplete				
Incomplete referral	referral				1
	Incomplete	Treatment			
Incomplete referral	referral	decision			1
	Communication				
	- patients /				
Incomplete referral	caregivers				1
Insufficient assessment	Transfer of				
(non-specific)	information				1
Insufficient assessment	Transfer of				
(non-specific)	patients				1
Insufficient assessment	Other				
(non-specific)					1
Insufficient assessment	Documentation	Documentation			
(non-specific)					1
Insufficient assessment	Documentation	Failure to identify			
(non-specific)		at risk/			
		vulnerable child			1
Insufficient assessment	Medication				
(non-specific)	prescribing				1
Insufficient assessment	Investigations				
(non-specific)					1
Insufficient assessment	Communication				
(non-specific)	- patients /				
	caregivers				1
Insufficient assessment	Communication				
(non-specific)	between				
	professionals				1
	Inadequate				
	discharge				
Investigations	planning				1
	Other				
Investigations	administrative				1

	Transfer of			
Investigations	information			1
	Investigations	Transfer of		
Investigations		patients		1
	Communication	Documentation		
	- patients /			
Investigations	caregivers			1
	Transfer of	Transfer of		
Investigations	information	information		1
Investigations	Documentation	Equipment		1
	Inadequate			
Medication	discharge			
administration	planning			1
Medication	Other			
administration	medication			1
Medication	Medication			
administration	administration			1
	Medication	Appointment	Transfer	
	monitoring	management	of	
Medication			informati	
administration			on	1
	Communication	Medication	Medicatio	
	- patients /	dispensing	n	
Medication	caregivers		prescribin	
administration			g	1
	Communication	Communication -		
Medication	- patients /	patients /		
administration	caregivers	caregivers		1
	Communication	Equipment		
Medication	- patients /			
administration	caregivers			1
	Equipment	Communication -		
Medication		patients /		
administration		caregivers		1
Medication	Documentation			
administration				1
	Medication	Communication -		
Medication	prescribing	patients /		
administration		caregivers		1

administration dispensing dispensing Documentation Communication - Medication patients / administration Access to care administration Medication Documentation Transfer of administration Documentation Medication Documentation administration Documentation Medication Documentation administration Documentation Medication Documentation Medication Medication Medication Medication Medication Medication Medication Medication		1 1 1
Medicationpatients /administrationAccess to careMedicationAccess to careadministrationDocumentationTransfer ofMedicationDocumentationMedicationDocumentationadministrationAccess to careMedicationInformationMedicationDocumentationAdministrationOtheradministrationMedication		1 1
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Medication administrationAccess to care DocumentationMedication administrationDocumentation informationMedication administrationDocumentation Other administrationMedication administrationOther medication		1 1
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Medication administration Documentation information Medication administration Documentation administration Medication administration Other administration		1
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Medication administration Documentation Medication administration Other administration		
administration Medication Other administration medication		
Medication Other administration medication		
administration medication		
Medication Medication		1
medication		
administration prescribing		1
Medication Equipment (Communi	
prescribing	cation -	
Medication p	patients /	
administration	caregivers	1
Medication Medication Medication		
administration dispensing prescribing		1
Communication		
Medication between		
administration professionals		1
Medication Equipment		
Medication dispensing dispensing		1
Medication dispensing Documentation		1
Other		
Medication dispensing medication		1
Medication Medication		
Medication dispensing prescribing prescribing		1
Medication dispensing Equipment		1
Other		
Medication dispensing administrative		1
Medication monitoring		1
Transfer of Transfer of		
Medication monitoring information information		1
Inadequate Transfer of		
Medication monitoring discharge information		1

	planning			
	Transfer of			
Medication monitoring	information			1
	Insufficient	Documentation		
	assessment			
Medication monitoring	(non-specific			1
Medication monitoring	Investigations			1
	Transfer of			
Medication prescribing	information			1
	Transfer of	Inadequate		
	information	discharge		
Medication prescribing		planning		1
Medication prescribing	Equipment			1
	Inadequate			
Medication prescribing	history			1
	Treatment	Communication		
	decision	between		
Medication prescribing		professionals		1
	Medication			
Medication prescribing	prescribing			1
	Medication	Access to care		
Medication prescribing	monitoring			1
	Communication			
	between			
Medication prescribing	professionals			1
	Medication	Medication		
Medication prescribing	dispensing	prescribing		1
	Inadequate	Documentation	Communi	
	history		cation -	
			patients /	
Medication prescribing			caregivers	1
	Communication			
	- patients /			
Medication prescribing	caregivers			1
	Medication			
Medication prescribing	dispensing			1
	Communication			
	- patients /			
Other	caregivers			1

Other caregivers professionals 1 Communication between Dother professionals 1 Other Other administrative		Communication	Communication			
Other professionals Other administrative Other Other Other Other Other Other When appropriate Other Other Other When appropriate Other Other Other Sailure to refer when appropriate Other Other Other Sailure to refer when appropriate Other Other Sailure to refer when appropriate Other Other Sailure to refer when appropriate Other Other Sailure to refer when administrative Other Sailure to refer administrative Other Sailure to refer when administrative Other Sailure to refer when appropriate Other Sailure to sidentify at risk/ Other Sailure to identify at risk/ Other Sailure to identify at risk/ Other Sailure to identify at risk/ Other Vullnerable child Inadequate discharge		- patients /	between			
Other professionals 1 Other Other administrative	Other	caregivers	professionals			1
Other professionals 1 Other administrative 1 Other Access to care Incomplete referral 1 Other Failure to refer when appropriate 1 Other Other Failure to refer when appropriate Referral administrative Other Other Other Inadequate triaging 1 Other Delayed referral Documentation identify at risk/vulnerable e child 1 Other Delayed referral Inadequate triaging 1 Other Failure to refer when appropriate 1 Other Failure to identify at risk/vulnerable child 1 Other Inadequate discharge 1		Communication				
Other administrative		between				
Other administrative 1 Access to care Incomplete referral 1 Other Other Failure to refer when appropriate Easilure to identify at risk with a	Other	professionals				1
Other		Other				
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Other Other when appropriate Failure to refer ent al managem administrative Other Other Other information administrative Delayed referral Documentation identify at risk/vulnerable triaging e history Other Delayed referral Documentation triaging e history Failure to refer when Appropriate Other appropriate Transfer of Other triaging e history Inadequate administrative Other appropriate Failure to refer when Appropriate Referral administrative Other issues Inadequate administrative Other issues Inadequate discharge		Access to care	Incomplete			
Other Other when appropriate 1 Failure to refer when appropriate ent al managem ent strativ e Other Other Other information Delayed referral Documentation entities 1 Other Delayed referral Documentation Failure to identify at risk/ vulnerable e history 1 Other Delayed referral Inadequate Inadequate triaging e history 1 Failure to refer when Other appropriate 1 Referral administrative 1 Referral administrative 1 Other Sissues 1 Failure to refer when 1 Referral administrative 1 Referral administrative 1 Inadequate Inadequate Inadequate Inadequate 1	Other		referral			1
Failure to refer when appropriate when appropriate when appropriate when appropriate when appropriate when appropriate ent at managem administrative e Other Other Other information administrative 1 Delayed referral Documentation identify at risk/ vulnerable of the child 1 Delayed referral Inadequate triaging e history 1 Failure to refer when appropriate e history 1 Referral administrative Other issues 1 Failure to refer when identify at risk/ vulnerable of identify at risk/ vulnerable child 1 Inadequate discharge			Failure to refer			
Other Other Other Other Irransfer of information administrative Other Other Irransfer of information administrative Irransfer of information administrative Irransfer of information administrative Irransfer of information administrative Irransfer of information Irransfer of	Other	Other	when appropriate			1
Other Other Other Other Other Information administrative Information Independent Pailure to refer when Other Appropriate Other Information Independent			Failure to refer	Appointm	Referr	
Other Other Other Other Other Other Other Other Information Administrative Information Other Information Informati			when appropriate	ent	al	
Other Other Other information administrative Inadequate administrative Other Information information administrative Inadequate Inadequate administrative Other Inadequate Inadeq				managem	admini	
Other Other Transfer of Other Information administrative Inadequate Administrative Inadequate Inadeministrative Inadequate Inadeministrative Inadeministrative Inadequate Inadeministrative Inad				ent	strativ	
Other information administrative 1 Delayed referral Documentation Failure to identify at risk/ vulnerabl e child 1 Other Delayed referral Inadequate triaging e history 1 Failure to refer when Other appropriate 1 Referral administrative Other issues 1 Failure to identify at risk/ vulnerable child 1 Inadequate discharge 1 Inadequate discharge 1 Inadequate 1 Inadequate discharge 1 Other information administrative 1 Other information administrative 1 Other vulnerable child 1 Inadequate discharge 1 Inadequate discharge 1 Inadequate 1 Inadequate Inadequate 1 Inadequate Inadequat					е	
Other information administrative 1 Delayed referral Documentation Failure to identify at risk/ vulnerabl e child 1 Other Delayed referral Inadequate triaging e history 1 Failure to refer when Appropriate 1 Referral administrative Other issues 1 Failure to identify at risk/ vulnerable child 1 Inadequate discharge 1 Other vulnerable child 1 Inadequate discharge 1 Inadequate discharge 1 Inadequate 1 Inadequate discharge 1 Inadequate 2 Inadequate 2 Inadequate 1 Inadequate 2 Inadequate 3 Inadequate 3 Inadequate 3 Inadequate 3 Inadequate 3 Inadequate 4 Inadequate 5 I	Other	Other			issues	1
Delayed referral Documentation Failure to identify at risk/ vulnerabl Polayed referral Delayed referral Inadequate Inadequat Pother Pailure to refer when Other appropriate Administrative Other issues 1 Failure to identify at risk/ Other vulnerable child Inadequate discharge		Transfer of	Other			
Other Delayed referral Inadequate triaging e history 1 Failure to refer when Other appropriate 1 Referral administrative Other issues 1 Failure to identify at risk/ Other vulnerable child Inadequate Inadequat e history 1 Referral administrative Inadequate Inadequat e history 1 Referral administrative Inadequate I	Other	information	administrative			1
Other Delayed referral Inadequate triaging e history 1 Failure to refer when Other appropriate Insurative Other issues 1 Failure to identify at risk/ Other vulnerable child Inadequate discharge		Delayed referral	Documentation	Failure to		
Other Delayed referral Inadequate triaging e history Tailure to refer when Other appropriate administrative Other issues Other railure to identify at risk/ vulnerable child Inadequate discharge				identify		
Other Delayed referral Inadequate Inadequat Other Triaging e history 1 Failure to refer When Other appropriate 1 Referral administrative Other issues 1 Failure to identify at risk/ Vulnerable child Inadequate discharge				at risk/		
Other Delayed referral Inadequate triaging e history 1 Failure to refer when Other appropriate 1 Referral administrative Other issues 1 Failure to identify at risk/ Other vulnerable child Inadequate discharge				vulnerabl		
Other triaging e history 1 Failure to refer when Other appropriate 1 Referral administrative Other issues 1 Failure to identify at risk/ Other vulnerable child Inadequate discharge	Other			e child		1
Failure to refer when Other appropriate 1 Referral administrative Other issues 1 Failure to identify at risk/ Vulnerable child 1 Inadequate discharge		Delayed referral	Inadequate	Inadequat		
Other appropriate 1 Referral administrative	Other		triaging	e history		1
Other appropriate 1 Referral administrative		Failure to refer				
Referral administrative Other issues 1 Failure to identify at risk/ Other vulnerable child 1 Inadequate discharge		when				
Other issues 1 Failure to identify at risk/ Other vulnerable child 1 Inadequate discharge	Other	appropriate				1
Other issues 1 Failure to identify at risk/ Other vulnerable child 1 Inadequate discharge		Referral				
Failure to identify at risk/ Other vulnerable child 1 Inadequate discharge		administrative				
identify at risk/ Vulnerable child Inadequate discharge	Other	issues				1
Other vulnerable child 1 Inadequate discharge		Failure to				
Inadequate discharge		identify at risk/				
discharge	Other	vulnerable child				1
		Inadequate				
Other planning 1		discharge				
	Other	planning				1

	Treatment and	Inadequate	Failure to		
	procedures	discharge	refer		
		planning	when		
			appropria		
Other			te		1
	Other	Other medication			
Other	medication				1
	Communication				
	- patients /				
Other administrative	caregivers				1
Other administrative	Appointment				
	management				1
Other administrative	Communication	Documentation			
	- patients /				
	caregivers				1
Other administrative	Communication				
	between				
	professionals				1
	Delayed referral	Inadequate	Failure to	Inadeq	
		triaging	identify	uate	
			at risk/	exami	
Other diagnosis and			vulnerabl	nation	
assessment			e child		1
Other medication	Medication	Medication			
	dispensing	prescribing			1
Referral administrative	Other				
issues	administrative				1
Referral administrative	Appointment				
issues	management				1
Referral administrative	Transfer of				
issues	information				1
	Communication				
Referral administrative	- patients /				
issues	caregivers				1
	Insufficient	Communication			
Referral administrative	assessment	between			
issues	(non-specific)	professionals			1
Referral administrative	Inadequate				
issues	triaging				1

	Inadequate			
	discharge			
Transfer of information	planning			1
	Inadequate	Communication -		
	discharge	patients /		
Transfer of information	planning	caregivers		1
Transfer of information	Access to care	Documentation		1
	Referral			
	administrative			
Transfer of information	issues			1
	Inadequate			
	discharge			
Transfer of information	planning			1
	Communication	Transfer of		
	between	information		
Transfer of information	professionals			1
	Inadequate			
Transfer of information	triaging			1
	Inadequate			
Transfer of information	history			1
	Access to care	Appointment		
Transfer of patients		management		1
	Transfer of			
Transfer of patients	information			1
	Incomplete	Incomplete		
Transfer of patients	referral	referral		1
	Failure to			
	identify at risk/			
Transfer of patients	vulnerable child			1
	Insufficient	Communication		
	assessment	between		
Transfer of patients	(non-specific)	professionals		1
	Delayed			
Transfer of patients	assessment			1
	Communication			
	- patients /			
Transfer of patients	caregivers			1
	Incomplete			
Transfer of patients	referral			1

	Communication			
	between			
Transfer of patients	professionals			1
	Diagnosis	Failure to identify	Investigat	
		at risk/	ions	
Transfer of patients		vulnerable child		1
	Treatment and	Communication -		
Treatment and	procedures	patients /		
procedures		caregivers		1
	Failure to refer			
Treatment and	when			
procedures	appropriate			1
Treatment and	Diagnosis			
procedures				1
	Insufficient			
Treatment and	assessment			
procedures	(non-specific)			1
	Other diagnosis	Failure to identify		
Treatment and	and assessment	at risk/		
procedures		vulnerable child		1
Treatment and	Treatment and	Treatment and		
procedures	procedures	procedures		1
Treatment and	Medication	Medication		
procedures	dispensing	dispensing		1
Treatment and	Investigations	Equipment		
procedures				1
Treatment and	Inadequate			
procedures	history			1
Treatment and	Appointment	Treatment and		
procedures	management	procedures		1
Treatment and	Access to care	Incomplete		
procedures		referral		1
	Transfer of	Failure to refer	Inadequat	
	information	when appropriate	е	
Treatment and			discharge	
procedures			planning	1
Treatment and	Transfer of			
procedures	patients			1
Treatment and	Delayed referral			1

procedures				
Treatment and	Incomplete			
procedures	referral			1
Treatment and	Incomplete	Incomplete		
procedures	referral	referral		1
	Referral	Appointment		
Treatment and	administrative	management		
procedures	issues			1
Treatment and	Diagnosis	Investigations	Investigat	
procedures			ions	1
Treatment and	Diagnosis	Investigations		
procedures				1
Treatment and	Inadequate	Inadequate		
procedures	triaging	history		1
Treatment and	Delayed	Transfer of		
procedures	assessment	information		1
	Delayed	Inadequate	Communi	
	assessment	triaging	cation	
			between	
Treatment and			profession	
procedures			als	1
	Delayed	Inadequate	Incomplet	
Treatment and	assessment	discharge	e referral	
procedures		planning		1
Treatment and	Treatment and	Appointment		
procedures	procedures	management		1
Treatment and	Medication			
procedures	dispensing			1
	Investigations	Transfer of	Inadequat	
		information	е	
Treatment and			discharge	
procedures			planning	1
	Investigations	Communication		
Treatment and		between		
procedures		professionals		1
Treatment and	Investigations	Equipment		
procedures				1
Treatment and	Investigations			
procedures				1

Treatment and	Equipment	Other			
procedures		administrative			1
Treatment decision	Access to care	Access to care			1
	Other diagnosis				
Treatment decision	and assessment				1
	Appointment				
Treatment decision	management				1
	Transfer of				
Treatment decision	information				1
	Diagnosis	Transfer of	Transfer		
		information	of		
			informati		
Treatment decision			on		1
	Insufficient				
	assessment				
Treatment decision	(non-specific)				1
	Inadequate	Inadequate	Document		
Treatment decision	triaging	history	ation		1
	Inadequate				
Treatment decision	examination				1
	Medication	Equipment			
Treatment decision	dispensing				1
	Other	Transfer of	Failure to		
	medication	information	identify		
			at risk/		
			vulnerabl		
Treatment decision			e child		1
	Other	Access to care	Document		
Treatment decision	medication		ation		1
Treatment decision	Equipment				1
Treatment decision	Delayed referral				1
	Transfer of	Incomplete	Delayed	Inadeq	
	information	referral	referral	uate	
				discha	
				rge	
				planni	
Treatment decision				ng	1
	Insufficient				
Treatment decision	assessment				1

	(non-specific			
	Inadequate			
Treatment decision	examination			1
	Failure to	Diagnosis		
	identify at risk/			
Treatment decision	vulnerable child			1
	Inadequate	Failure to identify		
	discharge	at risk/		
Treatment decision	planning	vulnerable child		1
	Other	Communication		
	medication	between		
Treatment decision		professionals		1
Treatment decision	Investigations			1
Treatment decision	Investigations	Documentation		1
	Investigations	Transfer of		
Treatment decision		information		1
	Communication	Communication -		
	between	patients /		
Treatment decision	professionals	caregivers		1
	Equipment	Transfer of		
Treatment decision		information		1
Treatment decision	Equipment	Investigations		1
	Equipment	Transfer of	Inadequat	
		information	е	
			discharge	
Treatment decision			planning	1
Grand Total				2191

3.4 The frequency of combinations of contributory factors for each primary incident type involving 'unwell' children

Contributory factors				Frequency of combination
Primary incident		T		
Medication dispensing				144
Treatment and procedures				87
Medication dispensing	Mistakes			78
Communication - patients / caregivers				76
Transfer of patient information				68
Inadequate triaging	Failure to			
	follow			
	protocol			67
Equipment				65
Inadequate triaging				63
Documentation				49
Delayed referral				46
Medication administering				38
Other				35
Investigations				34
Access to care				31
Treatment decisions				29
	Continuity of			
Transfer of patient information	care			27
Communication - patients / caregivers	Failure to			
	follow			26
	protocol			
Diagnostic issues				26
Medication dispensing	Mistake	Equipmen		
. 5		t/		
		medicatio		26
		n factors		
Delayed assessment				25
Transport/ transfer of patients				25
Inadequate triaging	Critical			
acquace c.iagiiig	thinking			22
Failure to refer when appropriate	Cillinaing			19
Medication prescribing				18

Other medication			18
Delayed referral	Failure to		
	follow		
	protocol		17
Incorrect/ incomplete referral			17
Inadequate triaging	Failure to	Critical	
	follow	thinking	
	protocol		15
Medication administering	Mistakes		14
Medication dispensing	Mistakes	Working	13
		conditions	13
Medication dispensing	Failure to		
	follow		12
	protocol		
Appointment management			10
	Failure to		
Delayed assessment	follow		10
	protocol		
Medication prescribing	Mistakes		10
Access to care	Continuity of		
	care		9
Deleverd	Continuity of		
Delayed assessment	care		9
Documentation	Failure to		
	follow		
	protocol		9
Inadequate discharge planning			8
Insufficient assessment (non-specific)			8
Medication administering	Patient age		8
Medication dispensing	Patient age		8
Medication prescribing	Patient age		8
Referral administrative issues			8
Treatment decisions	Working		
	conditions		8
Delayed referral	Critical		
	thinking		7
	Failure to		
	follow		
Failure to refer when appropriate	protocol		7

Medication dispensing	Equipment/			
	medication			7
	factors			
Medication dispensing	Inadequate			7
	guidelines			7
Medication dispensing	Mistakes	Failure to		
		follow		7
		protocol		
Other	Failure to			
	follow			
	protocol			7
Access to care	Failure to			
	follow			
	protocol			6
Communication - patients / caregivers	Mistakes			6
Delayed access out	Service			,
Delayed assessment	availability			6
Inadequate history taking				6
Inadequate history taking	Failure to			
	follow			
	protocol			6
Inadequate triaging	Patient/			
	caregiver			
	behaviour			6
Medication administering	Failure to			
	follow			6
	protocol			
Medication administering	Equipment/	Working		
	medication	conditions		6
	factors	Conditions		
Medication dispensing	Mistakes	Morking	Working	
		Working	conditio	6
		conditions	ns	
	Failure to			
	follow			
Transfer of patient information	protocol			6
Treatment and procedures	Patient/			
	caregiver			6
	behaviour			

Treatment and procedures	Continuity of			6
	care			
Communication between professionals				5
Delayed referral	Patient/			
	caregiver			
	behaviour			5
Delayed referral	Failure to	Critical		
	follow	thinking		
	protocol			5
Diagnostic issues	Failure to			
	follow			
	protocol			5
Other	Patient health			5
Other	Patient/			
	caregiver			
	behaviour			5
Other diagnosis and assessment				5
	Continuity of	Patient/		
	care	caregiver		
Transfer of patient information		geography		5
Treatment decisions	Inadequate			5
	guidelines			3
Access to care	Working			
	conditions			4
Communication between professionals	Failure to			
	follow			4
	protocol			
Communication - patients / caregivers	Patient age			4
Communication - patients / caregivers	Staff			
	knowledge			4
Communication - patients / caregivers	Failure to	Staff		
	follow	knowledg		4
	protocol	е		
	Inadequate			
Delayed assessment	guidelines			4
Deleved accessors	Working			4
Delayed assessment	conditions			4
Delayed referral	Staff			
	1	1	i I	

Inadequate triaging	Mistakes			4
Inadequate triaging	Education and			
	training			4
Medication administering	Patient/			
	caregiver			4
	behaviour			
Medication administering	Patient/			
	caregiver			4
	knowledge			
Medication dispensing	Working			4
	conditions			4
Medication dispensing	Mistake	Equipmen	Maylina	
		t/	Working conditio	4
		medicatio		4
		n factors	ns	
Medication prescribing	Failure to	Patient		
	follow	age		4
	protocol			
Treatment and procedures	Patient age			4
Treatment decisions	Failure to			
	follow			4
	protocol			
Treatment decisions	Continuity of			
	care			4
Access to care	Patient health			3
Access to care	Service			
	availability			3
Access to care	Patient/	Continuit		
	caregiver	y of care		
	geography			3
Communication between professionals	Continuity of			
	care			3
Communication between professionals	Working			_
·	conditions			3
Communication - patients / caregivers	Patient health			3
Communication - patients / caregivers	Patient/			
. 5	caregiver			3
	knowledge			
Communication - patients / caregivers	Inadequate			3
Exercise Parising and Parising				

	guidelines		
Communication - patients / caregivers	Continuity of		3
	care		3
Communication - patients / caregivers	Failure to	Patient	
	follow	age	3
	protocol		
Communication - patients / caregivers	Failure to	Critical	
	follow	thinking	3
	protocol		
Delayed referral	Continuity of		
	care		3
Delayed referral	Working		
	conditions		3
Delayed referral	Failure to	Staff	
	follow	knowledg	
	protocol	е	3
Diagnostic issues	Patient health		3
Diagnostic issues	Staff	Education	
	knowledge	and	
		training	3
Documentation	Mistakes		3
Equipment	Inadequate		3
	guidelines		
Equipment	Education and		3
	training		
Equipment	Working		3
	conditions		
Failure to identify at risk child			3
Inadequate examination			3
Inadequate triaging	Staff		
	knowledge		3
Inadequate triaging	Failure to	Patient	
	follow	age	
	protocol		3
Incorrect/ incomplete referral	Failure to		
	follow		
	protocol		3
Insufficient assessment (non-specific)	Patient age		3
Medication dispensing	Staff		3

	knowledge				
Medication dispensing	Mistakes	Patient			3
		age			3
Medication dispensing	Mistake	Inadequat			
		е			3
		guidelines			
Medication dispensing	Working	Working			3
	conditions	conditions			
Medication dispensing	Mistakes		Equipm		
		Working	ent/		3
		conditions	medicat		3
			ion		
Medication dispensing	Mistake	Equipmen	Working	Workin	
		t/	conditio	g	3
		medicatio		conditi	3
		n factors	ns	ons	
Medication prescribing	Staff				3
	knowledge				3
Other administrative					3
Other medication	Continuity of				3
	care				3
	Patient/				
	caregiver				
Transfer of patient information	geography				3
Treatment and procedures	Patient health				3
Treatment and procedures	Equipment/				
	medication				3
	factors				
Treatment and procedures	Working				3
	conditions				3
Communication between professionals	Mistake				2
Communication - patients / caregivers	Critical				2
	thinking				2
Communication - patients / caregivers	Education and				2
	training				2
Communication - patients / caregivers	Working				2
	conditions				
Communication - patients / caregivers	Failure to	Patient/			2
	follow	caregiver			2

	protocol	geography	
	Patient/		
Delayed assessment	caregiver		2
	geography		
	Failure to	Staff	
Delayed assessment	follow	knowledg	2
	protocol	е	
	Service	Inadequat	
Delayed assessment	availability	е	2
		guidelines	
	Service	Working	
Delayed assessment	availability	conditions	2
D. I. and	Service	Working	2
Delayed assessment	availability	conditions	2
Delayed referral	Mistakes		2
Delayed referral	Inadequate		
	guidelines		2
Documentation	Staff		
	knowledge		2
Equipment	Non-specific		
	organisational		2
	issues		
Failure to refer when appropriate	Patient age		2
	Staff		
Failure to refer when appropriate	knowledge		2
Inadequate discharge planning	Patient/		
	caregiver		
	behaviour		2
Inadequate discharge planning	Failure to		
	follow		
	protocol		2
Inadequate history taking	Critical		
	thinking		2
Inadequate triaging	Patient age		2
Inadequate triaging	Inadequate		
	guidelines		2
Inadequate triaging	Failure to	Patient	
	follow	health	
	protocol		2

Inadequate triaging	Critical	Patient	
	thinking	age	2
Inadequate triaging	Failure to	Staff	
	follow	knowledg	
	protocol	е	2
Inadequate triaging	Patient health	Critical	
		thinking	2
Inadequate triaging	Failure to	Education	
	follow	and	
	protocol	training	2
Incorrect/ incomplete referral	Patient age		2
Incorrect/ incomplete referral	Inadequate		
	guidelines		2
Incorrect/ incomplete referral	Education and		
	training		2
Incorrect/ incomplete referral	Failure to	Education	
	follow	and	
	protocol	training	2
Insufficient assessment (non-specific)	Failure to		
` '	follow		2
	protocol		
Insufficient assessment (non-specific)	Working		
, ,	conditions		2
Insufficient assessment (non-specific)	Patient age	Inadequat	
		е	2
		guidelines	
Investigations	Inadequate	3	
3	guidelines		2
Medication administering	Equipment/		
	medication		2
	factors		_
Medication administering	Education and		
medication administering	training		2
Medication administering	Continuity of		_
	care		2
Medication administering	Working		
5	conditions		2
Medication administering	Failure to	Patient/	1
medication administering	follow	caregiver	2
	Tottow	Calegivei	

	protocol	knowledg		
		е		
Medication dispensing	Education and			2
	training			2
Medication dispensing	Mistake	Staff		
		knowledg		2
		e		
Medication dispensing	Mistake	Mistake		2
Medication dispensing	Patient age	Mistake		2
Medication dispensing	Mistake	Other		
		staff		2
		factors		
Medication dispensing	Patient age	Working		2
		conditions		2
Medication dispensing	Staff	Education		
	knowledge	and		2
		training		
Medication dispensing	Mistakes	Equipmen	Failure	
		t/	to	
		medicatio	follow	2
		n factors	protoco	
			ι	
Medication dispensing	Mistake	Staff	Inadequ	
		knowledg	ate	2
			guidelin	2
		е	es	
Medication prescribing	Inadequate			2
	guidelines			2
Medication prescribing	Patient age	Failure to		
		follow		2
		protocol		
Medication prescribing	Mistakes	Equipmen		
		t/		2
		medicatio		
		n factors		
Other medication	Inadequate			2
other medication	guidelines			
	Patient/			
Transfer of patient information	caregiver			2

	behaviour		
Transport/ transfer of patients	Patient/		
	caregiver		
	geography		2
Transport/ transfer of patients	Failure to		
	follow		
	protocol		2
Transport/ transfer of patients	Service		
	availability		2
Transport/ transfer of patients	Working	Service	
	conditions	availabilit	
		у	2
Treatment and procedures	Mistakes		2
Treatment and procedures	Staff		
	knowledge		2
Treatment and procedures	Inadequate		
	guidelines		2
Treatment and procedures	Education and		
	training		2
Treatment and procedures	Service		
	availability		2
Treatment and procedures	Mistakes	Patient/	
		caregiver	2
		behaviour	
Treatment and procedures	Patient/	Patient	
	caregiver	health	2
	behaviour		
Treatment and procedures	Working	Continuit	2
	conditions	y of care	2
Treatment decisions	Equipment/		
	medication		2
	factors		
Treatment decisions	Continuity of	Inadequat	
	care	е	2
		guidelines	
Access to care	Patient/		
	caregiver		
	geography		1
Access to care	Patient/		1

	caregiver			
	knowledge			
Access to care	Patient/			
	caregiver			
	behaviour			1
Access to care	Critical			
	thinking			1
Access to care	Education and			
	training			1
Access to care	Continuity of	Patient		
	care	age		1
Access to care	Mistake	Failure to		
		follow		
		protocol		1
Access to care	Failure to	Staff		
	follow	knowledg		
	protocol	е		1
Access to care	Patient health	Failure to		
		follow		
		protocol		1
Access to care	Service	Continuit		
	availability	y of care		1
Access to care	Failure to	Service		
	follow	availabilit		
	protocol	у		1
Access to care	Looked-after	Patient/	Inadequ	
	child	caregiver	ate	
		geography	guidelin	
			es	1
Appointment management	Patient/			
	caregiver			
	geography			1
Appointment management	Patient age			1
Appointment management	Mistake			1
Appointment management	Service			
	availability			1
Appointment management	Working			
	conditions			1
Appointment management	Mistakes	Failure to		1

		follow		
		protocol		
Communication between professionals	Continuity of	Patient/		
	care	caregiver		1
		behaviour		
Communication between professionals	Failure to	Inadequat		
	follow	е		1
	protocol	guidelines		
Communication between professionals	Working	Continuit		1
	conditions	y of care		'
Communication between professionals	Critical	Failure to	Educati	
	thinking	follow	on and	1
		protocol	training	
Communication - patients / caregivers	Equipment/			
	medication			1
	factors			
Communication - patients / caregivers	Service			1
	availability			'
Communication - patients / caregivers	Patient/	Patient/		
	caregiver	caregiver		1
	knowledge	knowledg		'
		е		
Communication - patients / caregivers	Failure to	Patient		
	follow	health		1
	protocol			
Communication - patients / caregivers	Mistake	Patient		1
		age		'
Communication - patients / caregivers	Failure to	Working		
	follow	conditions		1
	protocol	condicions		
Communication - patients / caregivers	Failure to	Education		
	follow	and		1
	protocol	training		
Communication - patients / caregivers	Failure to	Staff	Patient	
	follow	knowledg	health	1
	protocol	е		
Communication - patients / caregivers	Patient	Service	Working	
	/caregiver	availabilit	conditio	1
	language	у	ns	

Follow protocol Follow pro	Communication - patients / caregivers	Failure to	Patient	Educati		
Communication - patients / caregivers follow protocol thinking available lity Communication - patients / caregivers Patient health Rowledge on and follow training protocol ol training availability caregiver geography service availability caregiver knowledg service availability of Patient Leave availability caregiver knowledg service availability of Patient health service availability of Patient health service availability follow protocol service availability follow service availability service availability follow service availability service availability		follow	age	on and		1
Follow protocol Patient Patien		protocol		training		
Patient Pati	Communication - patients / caregivers	Failure to	Critical	Service		
Communication - patients / caregivers and patients / caregivers are patients / caregivers and patients / caregivers are patients / caregiver and p		follow	thinking	availabi		1
Delayed assessment		protocol		lity		
Delayed assessment	Communication - patients / caregivers	Patient health			Failure	
Delayed assessment			Staff	Educati	to	
Delayed assessment			knowledg	on and	follow	1
Delayed assessment			е	training	protoc	
Delayed assessment Delayed assessment Delayed assessment Continuity of Patient/ care geography Service Patient/ caregiver geography Delayed assessment Delayed assessment Continuity of Patient/ caregiver knowledg e Education and Failure to Delayed assessment					ol	
Delayed assessment	Delayed accomment	Looked-after				1
Delayed assessment Knowledge Continuity of Patient / Care Caregiver geography Delayed assessment Care Caregiver Geography Caregiver Caregiver Caregiver Knowledg Early Caregiver Knowledg Early Caregiver Caregi	Detayed assessment	child				'
Continuity of Patient / Care Caregiver geography	Deleved	Staff				4
Delayed assessment Care geography Service Patient/ caregiver knowledg e Delayed assessment Continuity of Patient care health Delayed assessment Continuity of Continuity of Patient care health Delayed assessment Continuity of Failure to protocol Delayed assessment Continuity of Failure to protocol Service Failure to protocol Delayed assessment Education and Failure to protocol Delayed assessment Patient age Continuity of Follow protocol Continuity of Failure to protocol Continuity of Critical Delayed assessment Continuity of Continuity of Critical Delayed assessment Continuity of Continuity of Critical	Detayed assessment	knowledge				1
Delayed assessment Delayed assessment Delayed assessment Continuity of Patient knowledg e e Continuity of Patient health Delayed assessment Continuity of Patient age Continuity of Patient of Failure to protocol Delayed assessment Service Failure to protocol Delayed assessment Delayed assessment Education and training follow protocol Delayed assessment Delayed assessment Delayed assessment Delayed assessment Continuity of Failure to protocol Education and training follow protocol Delayed assessment Delayed assessment Continuity of Critical thinking Delayed assessment Continuity of Critical Torritical thinking Continuity of Critical Torritical thinking		Continuity of	Patient/			
Delayed assessment Service availability caregiver knowledg e Delayed assessment Continuity of care health Delayed assessment Continuity of care health Delayed assessment Continuity of care age Continuity of Failure to protocol Delayed assessment Service Failure to protocol Delayed assessment Education and training follow protocol Delayed assessment Delayed assessment Patient age Continuity of Critical thinking Continuity of Critical thinking Continuity of Critical thinking Continuity of Critical thinking	Delayed assessment	care	caregiver			1
Delayed assessment Delayed assessment Continuity of Patient care health Delayed assessment Continuity of Patient care health Delayed assessment Continuity of Patient age Continuity of Failure to protocol Delayed assessment Service Failure to protocol Delayed assessment Education and training follow protocol Delayed assessment Patient age Continuity of Critical thinking Continuity of Continuity of Critical thinking Continuity of Critical thinking 1			geography			
Delayed assessment Continuity of Patient health Delayed assessment Continuity of Patient care health Continuity of Patient care age Continuity of Failure to Delayed assessment Continuity of Failure to Delayed assessment Core follow protocol Service Failure to Delayed assessment Continuity of Failure to Delayed assessment Continuity of Patient to Delayed assessment Core follow protocol Failure to Delayed assessment Delayed assessment Continuity of Critical Thinking Continuity of Critical Toritical		Service	Patient/			
Delayed assessment Continuity of care health Delayed assessment Continuity of Patient care health Continuity of Patient care age Continuity of Failure to protocol Delayed assessment Service Failure to availability follow protocol Delayed assessment Education and Failure to protocol Delayed assessment Patient age Critical thinking Continuity of Critical Continuity of Patient age Critical thinking Continuity of Critical		availability	caregiver			
Delayed assessment Continuity of care health Delayed assessment Continuity of Patient care age Continuity of Failure to Delayed assessment Continuity of Failure to Continuity of Failure to Delayed assessment Service Failure to Delayed assessment Education and Failure to Delayed assessment Delayed assessment Education and Failure to Delayed assessment Patient age Critical thinking Continuity of Critical Continuity of Critical Taining follow	Delayed assessment		knowledg			1
Delayed assessment Care health Delayed assessment Continuity of care age Continuity of Failure to Delayed assessment Care follow 1 protocol Service Failure to availability follow 1 protocol Education and Failure to Delayed assessment Delayed assessment Training follow 1 Delayed assessment Patient age Critical thinking Delayed assessment Continuity of Critical 1			е			
Delayed assessment Continuity of Care age age age age age age age age age ag	Deleved	Continuity of	Patient			4
Delayed assessment Care age Continuity of Failure to Delayed assessment Care follow protocol Service Failure to Delayed assessment Delayed assessment Education and Failure to Delayed assessment Taining follow protocol Delayed assessment Delayed assessment Continuity of Critical Thinking Continuity of Critical Tailure to Thinking Continuity of Critical Thinking	Detayed assessment	care	health			1
Continuity of Failure to Delayed assessment Care follow protocol Service Failure to Delayed assessment Delayed assessment Education and Failure to Delayed assessment Training follow protocol Delayed assessment Patient age Critical thinking Delayed assessment Continuity of Critical Tollow protocol Thinking Tollow protocol	Deleved	Continuity of	Patient			4
Delayed assessment Care follow protocol Service Failure to availability follow protocol Education and Failure to Delayed assessment Delayed assessment Training follow protocol Patient age Critical thinking Delayed assessment Continuity of Critical 1	Detayed assessment	care	age			1
Delayed assessment Delayed assessment Delayed assessment Education and Failure to protocol Training follow protocol Delayed assessment Patient age Critical thinking Delayed assessment Continuity of Critical Telayed assessment		Continuity of	Failure to			
Delayed assessment Service Failure to availability follow protocol Education and Failure to training follow protocol Patient age Critical thinking Continuity of Critical Delayed assessment Service Failure to follow protocol Continuity of Critical thinking	Delayed assessment	care	follow			1
Delayed assessment availability follow protocol Education and Failure to training follow protocol Delayed assessment Patient age Critical thinking Delayed assessment Continuity of Critical 1			protocol			
protocol Education and Failure to training follow protocol Delayed assessment Patient age Critical thinking Delayed assessment Continuity of Critical 1		Service	Failure to			
Delayed assessment Education and Failure to training follow protocol Patient age Critical thinking Delayed assessment Continuity of Critical 1	Delayed assessment	availability	follow			1
Delayed assessment training follow protocol Delayed assessment Patient age Critical thinking Continuity of Critical 1			protocol			
Delayed assessment Patient age Critical thinking Continuity of Critical 1 Delayed assessment		Education and	Failure to			
Delayed assessment Patient age Critical thinking Continuity of Critical Delayed assessment 1	Delayed assessment	training	follow			1
Delayed assessment thinking 1 Continuity of Critical 1 Delayed assessment 1			protocol			
Delayed assessment Continuity of Critical 1		Patient age	Critical			_
Delayed assessment 1	Delayed assessment		thinking			1
Delayed assessment 1		Continuity of	Critical			
	Delayed assessment		thinking			1

	Patient	Service		
Delayed assessment	/caregiver	availabilit		1
	language	у		
	Patient/	Continuit		
Delayed assessment	caregiver	y of care		1
	geography			
	Patient health	Continuit		
Delayed assessment		y of care		1
Delevind assessment	Continuity of	Working		4
Delayed assessment	care	conditions		1
	Failure to	N/a white w		
Delayed assessment	follow	Working		1
	protocol	conditions		
	Working	Working		
Delayed assessment	conditions	conditions		1
	Critical	Education		
Delayed assessment	thinking	and		1
		training		
	Patient/	Service		
Delayed assessment	caregiver	availabilit		1
	behaviour	у		
	Failure to	Staff	Educati	
Delayed assessment	follow	knowledg	on and	1
	protocol	е	training	
Delayed referral	Patient			
	/caregiver			
	language			1
Delayed referral	Looked-after			
	child			1
Delayed referral	Education and			
	training			1
Delayed referral	Service			
	availability			1
Delayed referral	Patient/	Inadequat		
	caregiver	е		
	ethnicity	guidelines		1
Delayed referral	Failure to	Inadequat		
	follow	е		

Delayed referral	Mistakes	Inadequat			
		е			
		guidelines			1
Delayed referral	Patient	Service			
	/caregiver	availabilit			
	language	у			1
Delayed referral	Inadequate	Continuit			
	guidelines	y of care			1
Delayed referral	Working	Working			
	conditions	conditions			1
Delayed referral	Failure to	Education			
	follow	and			
	protocol	training			1
Delayed referral	Critical		Patient		
	thinking	Patient/	/		
		caregiver	caregiv		
		knowledg	er		
		е	geograp		
			hy		1
Delayed referral	Critical	Education	Patient		
	thinking	and			
		training	age		1
Delayed referral	Failure to	Critical	Staff		
	follow	thinking	knowle		
	protocol		dge		1
Delayed referral	Patient age	Failure to	Inadequ		
		follow	ate		
		protocol	guidelin		
			es		1
Delayed referral	Staff	Failure to	Working		
	knowledge	follow	conditio		
		protocol	ns		1
Delayed referral	Failure to	Staff	Educati		
	follow	knowledg	on and		
	protocol	е	training		1
Delayed referral	Service	Working	Working	Contin	
	availability	conditions	conditio	uity of	
		COMULIONS	ns	care	1

	caregiver		
	behaviour		
Diagnostic issues	Patient age		1
Diagnostic issues	Staff		
	knowledge		1
Diagnostic issues	Inadequate		
	guidelines		1
Diagnostic issues	Patient health	Staff	
		knowledg	
		е	1
Diagnostic issues	Failure to	Staff	
	follow	knowledg	
	protocol	e	1
Diagnostic issues	Inadequate	Staff	
	guidelines	knowledg	
		е	1
Diagnostic issues	Patient/	Working	
	caregiver	Working conditions	
	knowledge	Conditions	1
Documentation	Patient/		
	caregiver		
	knowledge		1
Documentation	Critical		
	thinking		1
Documentation	Continuity of		
	care		1
Documentation	Service		
	availability		1
Documentation	Failure to	Staff	
	follow	knowledg	
	protocol	е	1
Documentation	Failure to	Critical	
	follow	thinking	
	protocol		1
Documentation	Failure to		
	follow	Mistake	
	protocol		1
Documentation	Service	Continuit	
	availability	y of care	1

Documentation	Mistakes	Continuit		
		y of care		1
Equipment	Patient/			
	caregiver			1
	geography			
Equipment	Patient/			
	caregiver			1
	knowledge			
Equipment	Patient health			1
Equipment	Mistakes			1
Equipment	Equipment/			
	medication			1
	factors			
Equipment	Continuity of			1
	care			1
Equipment	Environmental			1
Equipment	Failure to	Inadequat		
	follow	е		1
	protocol	guidelines		
Equipment	Continuity of	Inadequat		
	care	е		1
		guidelines		
Equipment	Patient age	Continuit		1
		y of care		'
Equipment	Continuity of	Working		1
	care	conditions		1
Equipment	Staff	Education		
	knowledge	and		1
		training		
Equipment	Staff	Inadequat	Educati	
	knowledge	е	on and	1
		guidelines	training	
	Mistake	Continuit		
Failure to arrange follow up		y of care		1
Failure to identify at risk child	Critical			
	thinking			1
Failure to identify at risk child	Failure to	Critical		
	follow	thinking		
	protocol			1

Failure to identify at risk child	Inadequate	Education		
	guidelines	and		
		training		1
Failure to identify at risk child	Failure to	Staff	Educati	
	follow	knowledg	on and	
	protocol	е	training	1
	Patient			
	/caregiver			
Failure to refer when appropriate	language			1
	Critical			
Failure to refer when appropriate	thinking			1
	Education and	Patient/		
	training	caregiver		
Failure to refer when appropriate		behaviour		1
	Critical	Patient/		
	thinking	caregiver		
Failure to refer when appropriate		behaviour		1
	Failure to	Patient		
	follow	age		
Failure to refer when appropriate	protocol			1
	Failure to	Critical		
	follow	thinking		
Failure to refer when appropriate	protocol			1
	Continuity of	Critical		
Failure to refer when appropriate	care	thinking		1
	Failure to			
	follow	Mistake		
Failure to refer when appropriate	protocol			1
	Staff	Inadequat		
	knowledge	e		
Failure to refer when appropriate		guidelines		1
	Failure to	Continuit		
	follow	y of care		
Failure to refer when appropriate	protocol			1
	Service	Failure to	Patient	
	availability	follow	health	
Failure to refer when appropriate		protocol		1
	Critical	Failure to	Staff	
Failure to refer when appropriate	thinking	follow	knowle	1

		protocol	dge	
Inadequate discharge planning	Continuity of			
	care			1
Inadequate discharge planning	Service			
	availability			1
Inadequate discharge planning	Working			
	conditions			1
Inadequate discharge planning	Continuity of	Patient		
	care	health		1
Inadequate discharge planning	Patient age	Inadequat		
		е		
		guidelines		1
Inadequate discharge planning	Continuity of	Inadequat		
	care	е		1
		guidelines		
Inadequate discharge planning	Staff	Education		
	knowledge	and		
		training		1
Inadequate discharge planning	Staff	Education	Working	
	knowledge	and	conditio	
		training	ns	1
Inadequate examination	Patient/			
	caregiver			
	geography			1
Inadequate examination	Continuity of			
	care			1
Inadequate examination	Patient age	Failure to	Staff	
		follow	knowle	
		protocol	dge	1
Inadequate history taking	Patient/			
	caregiver			
	geography			1
Inadequate history taking	Service			
	availability			1
Inadequate history taking	Environmental			1
Inadequate history taking	Patient age	Patient		
		health		1
Inadequate history taking	Staff	Education		
	knowledge	and		1

		training		
Inadequate history taking	Failure to		Patient	
	follow		/	
	protocol	Working	caregiv	
		conditions	er	
			behavio	
			ur	1
Inadequate triaging	Patient health			1
Inadequate triaging	Patient/			
	caregiver			
	knowledge			1
Inadequate triaging	Equipment/			
	medication			
	factors			1
Inadequate triaging	Patient/	Patient/		
	caregiver	caregiver		
	behaviour	knowledg		
		е		1
Inadequate triaging	Failure to	Patient/		
	follow	caregiver		
	protocol	behaviour		1
Inadequate triaging	Critical	Patient		
	thinking	health		1
Inadequate triaging	Critical	Other		
	thinking	staff		
		factors		1
Inadequate triaging	Patient	Failure to		
	/caregiver	follow		
	language	protocol		1
Inadequate triaging	Patient health	Failure to		
		follow		
		protocol		1
Inadequate triaging	Failure to	Failure to		
	follow	follow		
	protocol	protocol		1
Inadequate triaging	Patient age	Failure to		
		follow		
		protocol		1
Inadequate triaging	Mistake	Failure to		1

		follow		
		protocol		
Inadequate triaging	Critical	Staff		
	thinking	knowledg		
		е		1
Inadequate triaging	Patient/	Critical		
	caregiver	thinking		
	behaviour			1
Inadequate triaging	Mistake	Critical		
		thinking		1
Inadequate triaging	Inadequate	Critical		
	guidelines	thinking		1
Inadequate triaging	Education and	Critical		
	training	thinking		1
Inadequate triaging	Service	Critical		
	availability	thinking		1
Inadequate triaging	Failure to			
	follow	Mistake		
	protocol			1
Inadequate triaging	Patient age	Inadequat		
		е		
		guidelines		1
Inadequate triaging	Critical	Inadequat		
	thinking	е		
		guidelines		1
Inadequate triaging	Failure to	Inadequat		
	follow	e		
	protocol	guidelines		1
Inadequate triaging	Inadequate	Continuit		
	guidelines	y of care		1
Inadequate triaging	Continuity of	Working		
	care	conditions		1
Inadequate triaging	Failure to	Service		
	follow	availabilit		
	protocol	у		1
Inadequate triaging	Inadequate	Service		
	guidelines	availabilit		
		у		1
Inadequate triaging	Failure to	Critical	Patient	1

	follow	thinking	/		
	protocol		caregiv		
			er		
			languag		
			e		
Inadequate triaging	Failure to	Critical	5		
	follow	thinking	Patient		
	protocol		age		1
Inadequate triaging	Patient/	Inadequat	Critical		
	caregiver	е	thinking		
	knowledge	guidelines			1
Inadequate triaging	Inadequate	Education	Critical		
	guidelines	and	thinking		
		training			1
Inadequate triaging	Failure to	Patient	Critical	Educat	
	follow	age	thinking	ion	
	protocol			and	
				trainin	
				g	1
Incorrect/ incomplete referral	Patient health			_	1
Incorrect/ incomplete referral	Critical				
	thinking				1
Incorrect/ incomplete referral	Staff				
	knowledge				1
Incorrect/ incomplete referral	Working				
	conditions				1
Incorrect/ incomplete referral	Mistakes	Patient			
		age			1
Incorrect/ incomplete referral	Continuity of	Failure to			
	care	follow			
		protocol			1
Incorrect/ incomplete referral	Patient age	Mistake			1
Incorrect/ incomplete referral	Inadequate	Education			
	guidelines	and			
		training			1
Incorrect/ incomplete referral	Patient/	Patient	Failure		
	caregiver	health	to		
	behaviour		follow		
			protoco		1
	1	l	l	l	

			l	
Insufficient assessment (non-specific)	Critical			
	thinking			1
Insufficient assessment (non-specific)	Mistake			1
Insufficient assessment (non-specific)	Continuity of			
	care			1
Insufficient assessment (non-specific)	Education and			
	training			1
Insufficient assessment (non-specific)	Continuity of	Patient/		
	care	caregiver		1
		geography		
Insufficient assessment (non-specific)	Failure to	Patient		
	follow	age		1
	protocol			
Insufficient assessment (non-specific)	Working	Working		
	conditions	conditions		1
Insufficient assessment (non-specific)	Failure to	Patient	Patient	
	follow	age	/	
	protocol		caregiv	1
			er	'
			knowle	
			dge	
Investigations	Patient health			1
Investigations	Patient/			
	caregiver			1
	behaviour			
Investigations	Failure to			
	follow			1
	protocol			
Investigations	Mistakes			1
Investigations	Equipment/			
	medication			1
	factors			
Investigations	Education and			1
	training			'
Investigations	Working			1
	conditions			
Investigations	Mistake	Patient/	Patient	4
		caregiver	,	1

Medication administering Inadequate guidelines Inadequate geography Inad			knowledg	caregiv	
Medication administering Inadequate guidelines 1 Medication administering Continuity of care caregiver geography 1 Medication administering Inadequate geography Patient/ caregiver behaviour 1 Medication administering Patient/ caregiver behaviour 1 Medication administering Patient age Patient/ caregiver knowledg e 1 Medication administering Patient age Patient/ caregiver knowledg e 1 Medication administering Patient/ caregiver knowledg e 1 Medication administering Mistakes Patient/ caregiver knowledg e 1 Medication administering Staff patient (caregiver knowledge) 1 Medication administering Staff patient (caregiver knowledge) 1 Medication administering Patient (caregiver knowledge) 1 Medication administering<			е	er	
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follow protocol Medication administering Patient age Mistake 1	Medication administering	Mistake			
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Medication administering Patient age Mistake 1			protocol		
	Medication administering	Patient age			1
	Medication administering	Failure to	Failure to		1

	follow	follow		
	protocol	protocol		
Medication administering	Mistakes	Equipmen		
		t/		
		medicatio		1
		n factors		
Medication administering	Mistake	Equipmen		
		t/		
		medicatio		1
		n factors		
Medication administering	Inadequate	Continuit		
	guidelines	y of care		1
Medication administering	Mistake	Working		1
	mistane	conditions		
Medication administering	Mistakes	Working		1
		conditions		'
Medication administering	Mistake	Working		1
		conditions		'
Medication administering	Mistake	Working		1
		conditions		'
Medication administering	Working	Working		4
	conditions	conditions		'
Medication administering	Inadequate	Education		
	guidelines	and		1
		training		
Medication administering	Patient/	Education		
	caregiver	and		1
	knowledge	training		
Medication administering	Patient age		Staff	
		Mistake	knowle	1
			dge	
Medication administering	Patient health	Patient/	Continu	
		caregiver	ity of	4
		knowledg	care	1
		е		
Medication administering	Mistake	Failure to	Continu	
		follow	ity of	1
		protocol	care	
Medication administering	Failure to	Working	Working	1

	follow	conditions	conditio		
	protocol		ns		
Medication administering	Mistake	Working	Working		
		conditions	conditio		1
			ns		
Medication administering	Mistakes	Failure to	Working		
		follow	conditio		1
		protocol	ns		
Medication administering	Patient age	Working	Working		
		conditions	conditio		1
		Conditions	ns		
Medication administering	Mistake	Working	Educati		
		conditions	on and		1
		Conditions	training		
Medication administering	Mistake	Patient	Failure	F	
		health	to	Equip	
			follow	ment/	1
			protoco	medic	
			ι	ation	
Medication administering	Continuity of	Patient/	Patient		
	care	caregiver	/	Inadeq	
		geography	caregiv	uate	
			er	protoc	1
			knowle	ols	
			dge		
Medication dispensing	Patient health				1
Medication dispensing	Patient/				
	caregiver				1
	behaviour				
Medication dispensing	Failure to	Patient			
	follow	health			1
	protocol				
Medication dispensing	Patient age	Failure to			
-		follow			1
		protocol			
Medication dispensing	Working	Staff			
	conditions	knowledg			1
		e			
Medication dispensing	Equipment/	Mistake			1
	_4=.6				•

	medication			
	factors			
Medication dispensing	Staff	Equipmen		
	knowledge	t/		4
		medicatio		1
		n factors		
Medication dispensing	Mistake	Service		
		availabilit		1
		у		
Medication dispensing	Working	Continuit		
	conditions	y of care		1
Medication dispensing	Mistakes	Working		
		conditions		1
Medication dispensing	Staff	Working		
3	knowledge	conditions		1
Medication dispensing	Failure to	Education		
meaneacon and ponoming	follow	and		1
	protocol	training		
Medication dispensing	Mistake	Education		
Medication dispensing	Mistake	and		1
		training		'
Medication dispensing	Inadequate	Education		
medication dispensing	guidelines	and		1
	guidetilles	training		1
Madiantian dispansing	Dationton	Failure to		
Medication dispensing	Patient age		Mistalia	4
		follow	Mistake	1
AA IP A I I I I I I I I I I I I I I I I	Add to Land	protocol		
Medication dispensing	Mistake	Equipmen		
		t/	Mistake	1
		medicatio		
		n factors		
Medication dispensing	Mistakes	Working	Other	
		conditions	staff	1
			factors	
Medication dispensing	Mistake	Patient	Equipm	
		age	ent/	1
			medicat	'
			ion	
Medication dispensing	Patient health	Mistake	Equipm	1

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			ent/ medicat	
			ion	
Medication dispensing	Mistakes	Patient	Inadequ	
3		age	ate	
			guidelin	1
			es	
Medication dispensing	Equipment/	Failure to	Inadequ	
eareacien alapenanig	medication	follow	ate	
	factors	protocol	guidelin	1
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Medication dispensing	Failure to		Inadequ	
medication disperising	follow	Staff	ate	
	protocol	knowledg	guidelin	1
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Addination dispension	Mintale	Education	es Marking	
Medication dispensing	Mistake		Working	4
		and	conditio	1
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Medication dispensing	Patient age	Failure to	Working	
		follow	conditio	1
		protocol	ns	
Medication dispensing	Mistake	Failure to	Working	
		follow	conditio	1
		protocol	ns	
Medication dispensing	Mistake	Staff	Working	
		knowledg	conditio	1
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Medication dispensing	Mistake	Working	Working	
		conditions	conditio	1
		Conditions	ns	
Medication dispensing	Mistake		Working	
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Medication dispensing	Patient age	Staff	Educati	
		knowledg	on and	1
		е	training	
Medication dispensing	Failure to	Equipmen	Educati	
	follow	t/	on and	1
	protocol	medicatio	training	
	-			

		n factors			
Medication dispensing	Inadequate	Continuit	Patient	Patien	
	guidelines	y of care	/	t/	
			caregiv	caregi	4
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			dge	our	
Medication dispensing	Failure to	\\\	Staff	Datian	
	follow	Working	knowle	Patien	1
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Medication dispensing	Mistake		Working	Workin	
		Mistake	_	g	4
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Medication dispensing	Mistakes	Patient	NA/a włożna w	Workin	
		age	Working	g	_
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			ns	ons	
Medication prescribing	Patient/				
	caregiver				1
	geography				
Medication prescribing	Patient				
	/caregiver				1
	language				
Medication prescribing	Patient/				
	caregiver				1
	behaviour				
Medication prescribing	Patient health				1
Medication prescribing	Failure to				
	follow				1
	protocol				
Medication prescribing	Patient/	Patient/			
	caregiver	caregiver			4
	geography	knowledg			1
		е			
Medication prescribing	Patient/	Patient/			
	caregiver	caregiver			_
	behaviour	knowledg			1
		е			

Medication prescribing	Mistake	Patient			1
		age			'
Medication prescribing	Patient/	Failure to			
	caregiver	follow			1
	behaviour	protocol			
Medication prescribing	Mistake	Failure to			
		follow			1
		protocol			
Medication prescribing	Mistake	Staff			
		knowledg			1
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Medication prescribing	Patient health	Mistake			1
Medication prescribing	Patient age	Equipmen			
		t/			1
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		n factors			
Medication prescribing	Failure to	Inadequat			
	follow	e			1
	protocol	guidelines			
Medication prescribing	Continuity of	Inadequat			
	care	е			1
		guidelines			
Medication prescribing	Staff	Failure to	Patient		
	knowledge	follow			1
		protocol	age		
Medication prescribing	Mistakes	Staff	Dationt		
		knowledg	Patient		1
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		knowledg			1
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Medication prescribing	Continuity of	Inadequat	Dationt	1	
	care	е	Patient		1
		guidelines	age		
Other	Patient/				
	caregiver				
	geography				1
Other	Looked-after				
	child				1

Other	Inadequate			
	guidelines			1
Other	Continuity of			
	care			1
Other	Working			
	conditions			1
Other	Continuity of	Patient/		
	care	caregiver		
		geography		1
Other	Continuity of	Patient		
	care	health		1
Other	Inadequate	Looked-		
	guidelines	after		
		child		1
Other	Continuity of	Patient		
	care	age		1
Other	Continuity of	Failure to		
	care	follow		
		protocol		1
Other	Continuity of	Inadequat		
	care	e		
		guidelines		1
Other	Staff	Working		
	knowledge	conditions		1
Other	Failure to	Staff	Educati	
	follow	knowledg	on and	
	protocol	е	training	1
Other administrative	Patient age			1
Other administrative	Failure to			
	follow			
	protocol			1
Other administrative	Mistake			1
Other administrative	Inadequate			
	guidelines			 1
	Patient age	Staff		
		knowledg		
Other diagnosis and assessment		e		 1
Other medication	Patient/			 1
	caregiver			

	geography			
Other medication	Patient/			
	caregiver			1
	behaviour			
Other medication	Mistakes			1
Other medication	Education and			
	training			1
Other medication	Service	Patient/		
	availability	caregiver		1
		behaviour		
Other medication	Mistake	Staff		
	mstarte	knowledg		1
		e		'
Other medication	Mistakes	Staff	Other	
Other medication	Mistakes			4
		knowledg	staff	'
		е	factors	
Other medication	Mistake	Staff	Educati	
		knowledg	on and	1
		е	training	
Referral administrative issues	Mistake			1
Referral administrative issues	Service			
	availability			1
Referral administrative issues	Failure to	Patient		
	follow	health		
	protocol			1
Referral administrative issues	Continuity of	Patient/		
	care	caregiver		
		knowledg		
		e		1
Referral administrative issues	Continuity of	Inadequat		
	care	е		
		guidelines		1
Referral administrative issues	Education and	Working		
	training	conditions		1
Referral administrative issues	Looked-after	Patient/	Continu	
	child	caregiver	ity of	
		geography	care	1
	Patient	22		
Transfer of patient information	/caregiver			1
ransier of patient information	/ Curegiver			

	language			
Transfer of patient information	Mistakes			1
	Working			
Transfer of patient information	conditions			1
	Failure to	Patient		
	follow	health		
Transfer of patient information	protocol			1
	Continuity of	Patient		
Transfer of patient information	care	health		1
	Failure to	Critical		
	follow	thinking		
Transfer of patient information	protocol			1
	Mistakes	Equipmen		
		t/		
		medicatio		
Transfer of patient information		n factors		1
	Looked-after	Continuit		
Transfer of patient information	child	y of care		1
	Continuity of	Continuit		
Transfer of patient information	care	y of care		1
	Failure to	Continuit		
	follow	y of care		
Transfer of patient information	protocol			1
	Working	Working		
Transfer of patient information	conditions	conditions		1
	Continuity of	Education		
	care	and		
Transfer of patient information		training		1
	Failure to	Failure to	Working	
	follow	follow	conditio	
Transfer of patient information	protocol	protocol	ns	1
Transport/ transfer of patients	Patient/			
	caregiver			
	behaviour			1
Transport/ transfer of patients	Staff			
	knowledge			1
Transport/ transfer of patients	Inadequate			
	guidelines			1
Transport/ transfer of patients	Environmental			1

Transport/ transfer of patients	Patient health	Staff			
		knowledg			
		е			1
Transport/ transfer of patients	Service	Working			
	availability	conditions			1
Transport/ transfer of patients	Failure to	Education			
	follow	and			
	protocol	training			1
Transport/ transfer of patients	Failure to	Service	Continu		
	follow	availabilit	ity of		
	protocol	у	care		1
Transport/ transfer of patients	Service		Working		
	availability	Working	conditio		
		conditions	ns		1
Transport/ transfer of patients	Staff	Patient	Inadequ	Educat	
·	knowledge	age	ate	ion	
			guidelin	and	
			es	trainin	
				g	1
Treatment and procedures	Patient			3	
·	/caregiver				1
	language				
Treatment and procedures	Critical	Patient			
·	thinking	health			1
Treatment and procedures	Continuity of	Patient/			
	care	caregiver			
		knowledg			1
		e			
Treatment and procedures	Inadequate	Staff			
·	guidelines	knowledg			1
	3	e			
Treatment and procedures	Environmental	Inadequat			
		е			1
		guidelines			
Treatment and procedures	Education and	Working			
, and p. 55552.65	training	conditions			1
Treatment and procedures	Patient health	Working			
	. deserte medicin	conditions			1
Treatment and procedures	Failure to	Working			1
Treatment and procedures	i altule to	MOLKING			1

	follow	conditions			
	protocol				
Treatment and procedures	Equipment/ medication	Working conditions			1
-	factors	N/ 1:			
Treatment and procedures	Working	Working conditions			1
Treatment and procedures	conditions Mistake	Working			
Treatment and procedures	MISLAKE	conditions			1
Treatment and procedures	Patient age	Education			
rreatment and procedures	Patient age	and			1
		training			1
Treatment and procedures	Equipment/	Education			
Treatment and procedures	medication	and			1
	factors	training			•
Treatment and procedures	Continuity of	Looked-	Patient		
Treatment and procedures	care	after	/		
		child	caregiv		
			er		1
			geograp		
			hy		
Treatment and procedures	Continuity of	Failure to	Patient		
	care	follow	health		1
		protocol			
Treatment and procedures	Mistake	N/a white w	Working		
		Working conditions	conditio		1
		Conditions	ns		
Treatment and procedures	Failure to	Working	Educati		
	follow	conditions	on and		1
	protocol	condicions	training		
Treatment and procedures	Patient age	Patient/	Educati	Contin	
		caregiver	on and	uity of	1
		geography	training	care	
Treatment decisions	Patient age				1
Treatment decisions	Staff				1
	knowledge				·
Treatment decisions	Mistake				1
Treatment decisions	Failure to	Failure to			1
	follow	follow			

	protocol	protocol		
Treatment decisions	Patient age	Inadequat		
		е		1
		guidelines		
Treatment decisions	Patient/	Inadequat		
	caregiver	е		1
	behaviour	guidelines		
Treatment decisions	Staff	Inadequat		
	knowledge	е		1
		guidelines		
Treatment decisions	Patient age	Continuit		1
		y of care		'
Treatment decisions	Inadequate	Continuit		1
	guidelines	y of care		'
Treatment decisions	Staff	Continuit		1
	knowledge	y of care		'
Treatment decisions	Failure to	Staff	Patient	
	follow	knowledg		1
	protocol	е	age	
Treatment decisions	Continuity of	Inadequat	Working	
	care	е	conditio	1
		guidelines	ns	
Grand Total				2191

3.5 Medline ovid search strategy for literature review

- 1. (Vaccin* adj3 error*).mp.
- 2. (Vaccin* adj3 program* error*).mp
- 3. (Immuni* adj3 error*).mp.
- 4. Immuni* safety.mp.
- 5. Vaccinat* safety.mp.
- 6. Medicat* error*.mp.
- 7. (triag* adj3 error*).
- 8. (triag* adj3 incident*).mp.
- 9. (clinical assessment adj3 error*).mp.
- 10. (patient assessment adj3 error*).mp.
- 11. (assessment adj3 safety incident).mp.
- 12. "diagnos* error*".mp.
- 13. diagnos* incident*.mp.
- 14. (patient* record* adj3 error*).mp.
- 15. (medical record* adj3 error*).mp.
- 16. (referral* adj3 error*).mp.
- 17. (referral* adj3 safety).mp.
- 18. (referral* adj3 incident*).mp.
- 19. (communicat* adj3 error*).mp.
- 20. (communicat* adj3 failure).mp.
- 21. communicat* incident*.mp.
- 22. (communicat* adj3 patient* safety).mp.
- 23. (1-22)/or
- 24. exp Child Health Services/ or exp Child, Preschool/ or exp Child/
- 25. Paediatri*.mp.
- 26. Pediatri*.mp.
- 27. exp Adolescent/ or exp Adolescent Health Services/
- 28. exp Infant/
- 29. exp Infant, Newborn/
- 30. (23-28)/or
- 31. (Improve* adj3 interven*).mp.
- 32. exp Quality Improvement/
- 33. (error* adj3 prevent*).mp.
- 34. (safety adj3 improve*).mp.
- 35. (error* adj3 reduc*).mp.
- 36. (31-35)/or
- 37. Animals/
- 38. animal stud*.mp.
- 39. 37 or 38
- 40. 23 AND 30 AND 36
- 41. 40 NOT 39