PAIN



The establishment, maintenance, and adaptation of high- and low-impact chronic pain: a framework for biopsychosocial pain research

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1. Introduction

We present a framework for the study of states of chronic pain and transitions between those states. We capture in the framework the dynamic nature of pain: people live with pain that changes over time. First, we offer definitions of both acute and chronic pain and explore the contextual considerations related to the common use of this temporal dichotomy. Second, we promote the importance of incorporating the impact pain has on a person's life. Finally, we discuss the challenges and opportunities inherent in implementing this common approach. Our goal is to produce a framework for the study of the development, maintenance, and resolution of chronic pain.

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Whether a single brief event or a constant feature of life, pain interrupts to prioritise protection, interferes with activity, reduces quality of life, and can alter identity.⁴⁴ Protection is achieved by escape from harm, avoidance of perceived danger, withdrawal for respite and repair, and communication of incapacity and environmental risk; longer-term protection is achieved by learning the cues for pain and injury.⁵³ From this perspective, pain is most usefully considered a need state, fundamentally a motivational drive to protect.⁴⁹ This approach centres our attention on the consequences of pain for the person in their context, on its duration and its impact.

2. A person's pain status

Pain is defined as "an unpleasant sensory and emotional experience associated with, or resembling that associated with, actual or potential tissue damage."³⁴ There is a logical case of a state of "no pain," but to have no pain is a rare occurrence, only recently made possible by the advent of anaesthesia/analgesia. A state of the continuous absence of any pain at all is profoundly abnormal, appearing only as congenital nociceptor deficiency or dysfunction. It is far from adaptive, notably leading to major clinical problems associated with the absence of defensive responding and learning with a consequent severe shortening of life expectancy.^{9,50,51}

2.1. Acute and chronic

The terms "acute" and "chronic" represent a temporal dichotomy, with "acute" meaning short lived or immediate, and chronic meaning long term. Colloquially, however, both can be used interchangeably to mean "bad," causing confusion in clinical encounters when patients use these as terms for severity or impact and clinicians use them for temporality.⁵

In this article, we use "acute pain" to mean pain of short duration, without any implication of severity or urgency. The pragmatic challenge in its use is the length of this short duration. Acute pain is typically defined as lasting from onset to 3 months in duration and so encompasses everything from a momentary muscular "cramp" to postoperative pain.^{27,52} Therefore, acute pain is both common and a normal part of everyday life. The few studies that establish the base rate of everyday pain generally report high incidence. For example, in 1 observational study of everyday pain in 3- to 7-year-olds, the event rate was 0.33 incidents per hour per child.³¹ Painful bumps and

scrapes in the playground are normal. For adults, episodes of naturally occurring acute pain are also common. In a Europeanwide study of 8506 patients, 70% of adults reported at least 1 pain event a month (such as headache, menstrual pain, and muscle pain).⁴⁸ Acute pain can also occur deliberately as part of a social process (eg, body adomment, contact sport, or ritual). However, most everyday acute pain does not require clinical intervention; it is self-limiting.

Clinical studies often focus on pain related to medical procedures or that occurs as the result of illness, disease, or injury (accidental, self-induced, or medically induced), and although acute pain can be relatively straightforward to manage, in particular when the timing and extent of trauma are controlled, it is not always so simple. Acute pain can be complicated for healthcare professionals to assess and manage when there are concurrent symptoms or conditions, it occurs in the presence of chronic pain, or when communication about pain is difficult, for instance in an emergency.²³

Furthermore, we recognise that the 3-month limit is an arbitrary distinction that can lead to problems. For example, it does not account for life stage. Consider the case of new-born babies who receive repeated needle sticks for diagnostic tests: Each pain may be acute, but the pain has been present for most of the child's life. For a baby, less than 3 months is not a "short" duration. These and other concerns have led some to question the focus on duration in definitions, suggesting instead a focus on presumed mechanism.⁸

Pain of longer duration is known as "chronic pain." Typically, chronic means having lasted for 3 months or more, as captured in the IASP definition and more recently included in *ICD-11.*³⁸ Historically, chronic pain was considered to start at 6 months for adults and 3 months for children. The problem of definitions that privilege duration has been discussed,⁴⁶ and 3 months for adult and child is now thought more clinically relevant. Given the consensus on the use of a 3-month definition, this serves as the dividing boundary that differentiates chronic from acute pain.

This definition has some difficulties. As with acute pain, 3 months is an absolute cutoff: 3 months for a 6-month-old is 50% of a life, whereas for a 60-year-old, 3 months is 0.4% of a life. If a child of 2 months has lived their life in pain, this definition of chronic would not apply, which in some circumstances would be more clinically meaningful.²⁵ Furthermore, many people report pain that changes in quality or location and which fluctuates and/ or is episodic. Consider that the International Headache Society classifies a number of chronic headaches as episodic with different decisions about the number, frequency, and extent of episodes,²⁰ as do the Rome IV Diagnostic Criteria for Functional Gastrointestinal Disorders, which are less exact and use phrases such as "continuous," "nearly continuous," and "intermittent."37 Including frequency in the definition of chronic low back pain may be useful and lead to a more accurate identification of treatment responders.²² In short, a category of chronic pain needs to encompass the experience of people who have intermittent, episodic, or continuous pain, pain of different quality and intensity, and the pain may be a sole primary complaint, secondary to disease and illness, or one of several chronic complaints.^{29,43}

The variability in nosology emerging from different clinical specialities, and considerations of duration in the context of longevity, does not negate the value of a simple duration dichotomy (acute-chronic). We argue for the informed use of context when considering its use, and the need to look beyond simple labels when combining data or insights. It is a useful starting point to explore the specific features of that temporal definition in context.

2.2. High- and low-impact acute and chronic pain

Pain can impact multiple aspects of a person's life. We prefer "impact" over other common terms, such as disability, suffering, or distress, because it draws attention to the diverse effects of pain on the person. It is generally used conditionally to refer to the consequences of pain on a particular outcome.^{13,39,42}

Duration is not a good predictor of impact. A momentary pain of an accident, incident, or procedure can have drastic effects on a person. One example is the cumulative effects of repeated exposure to acute pain in new-born babies on brain development.² Another is the potential role of discrete painful events as psychologically traumatic, leading to major impact.³⁵ And, a third is of uncontrolled pain near end of life, which may last less than 3 months but which can be devastating for a person and significant others.²⁸

Discriminating by impact is more common in considerations of chronic pain. In particular, "high-impact chronic pain"has been defined in the United States as activity limitation⁴⁷ and later as activity and participation limitation,^{33,39} contrasted with a category of chronic pain without limitations. This thinking was more recently captured in the Graded Chronic Pain Scale Revised as high impact compared with mild or bothersome pain.⁴⁵ These categories allow for greater discrimination when trying to bridge between population-based studies of prevalence of chronic pain and clinical studies with adults expressing healthcare needs. The prevalence of adults with high-impact chronic pain is more typically estimated conservatively as at least 5%, in contrast to the headline population figures for all chronic pain, conservatively estimated at 20%.^{33,55}

The idea that chronic pain can have low or no impact is an interesting one. Indeed, the potential for the existence of pain without impact is at the heart of the biopsychosocial model^{6,16,17} and a treatment goal in psychological rehabilitation.⁵⁴ Although the complete resolution of chronic pain is desirable as a treatment objective, the transition to a state of low(er) impact chronic pain is often more realistic and still an important objective for individuals, healthcare providers, and society. An example is in the context of normal ageing with accommodation to life with increasingly unreachable goals achieved by altering those goals.¹²

For our purposes, high impact is defined by the extent of difficulties in function and disability (self-care, occupational engagement, and social activities)²⁶ in line with the WHO. Again, we propose an informed and context-dependent use, with the need to look beyond simple labels when combining data or insights. It is a useful starting point to explore the specific features of how high impact can be determined from the available measurement.

3. States and transition

Taking duration and impact together, we propose a transitional framework for the study of 5 categorical "states" (**Table 1** and **Fig. 1**), which include acute low-impact pain, acute high-impact pain, chronic low-impact pain, chronic high-impact pain, and a "resolved" no chronic pain state.

Although we refer here to pain duration (acute, chronic) and impact (high, low) as dichotomies, we recognise the continuous, overlapping and dynamic aspect of the pain experience. For many people, pain is an additional burden to other diseases. Our choices here are illustrative not ontological providing a framework for investigation—placing an emphasis on measurement and its use within individual investigations. In line with the US pain strategy,²⁶ we recognise that introducing categories creates

States of pain.			
Duration	Impact	Features	
Acute pain	Low impact	Pain of less than 3-mo duration that is not associated with major self-care, occupational, and social activity restrictions Pain of less than 3-mo duration can occur in a normal everyday context. Examples are inoculation, minor injury, or pain incidental to aesthetics or recreation	
Acute pain	High impact	Pain of less than 3-mo duration that is associated with major social, personal, or role restrictions Pain of less than 3-mo duration can have a major impact. Examples are major trauma, headache, end of life pain	
Chronic pain	Low impact	Pain of 3 mo or more duration that is not associated with major social, personal or role restrictions Pain of 3 mo or longer can be related to a disease or be a primary disorder of the nervous system but can have minor impact	
Chronic pain	High impact	Pain of 3 mo or more duration that is associated with major social, personal, or role restrictions Pain of 3 mo or longer can be related to a disease or be a primary disorder of the nervous system but can have major impact	
Chronic pain (resolution)	No impact	The chronic pain of interest can progress to a new state of resolved chronic pain This new state is similar to acute pain of low impact in which everyday pain may occur but is always in the context of having had chronic pain	

opportunities for research, in particular population-based research, but can under some circumstances lead to a statistical loss of information.

Table 2 outlines 10 possible trajectories of change in states, representing transitions (or absence of transition) in a person's pain state. We are interested in the onset of chronic pain, whether it is low or high impact, and its starting point of low- or high-impact acute pain. We are also interested in no change, or the maintenance of chronic pain, whether low or high impact, and the factors that lead to people becoming "stuck" in their pain state.³ And finally, we are interested in change in state, worsening from low impact to high impact, or improving from high impact to low impact, or a resolution from chronic low- or high-impact pain back: transition to a new normal state in which the specific pain(s) meeting the criteria for chronicity has/have resolved, but the

natural rate of everyday pain resumes. These states and transitions are outlined in **Figure 1**.

4. Further considerations

Our focus on duration and impact raises several issues for consideration:

- (1) Pain can be described by its pathological cause, mechanism, intensity, location, frequency, diurnality, or as a collection of features in a measure of severity. Such features are important, but, in this framework, they would be held in analyses as potential predictors, correlates, or process variables in an examination of impact and duration rather than part of their definition.
- (2) The premise that a person can have chronic pain with low impact clashes with the *ICD-11* definition of primary chronic

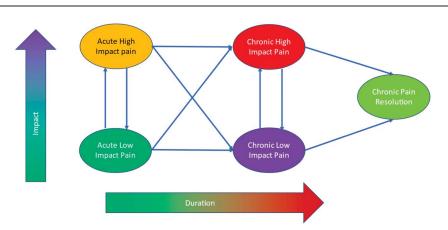


Figure 1. A framework for the establishment, maintenance, and adaptation of high- and low-impact chronic pain: Pain is described with 2 dominant dimensions: duration and impact. Both are dichotomised to establish specific states. Duration is split into acute pain defined as pain lasting less than 3 months and chronic as pain of 3-month duration or longer. Impact is split into low and high impact defined as a minor or major impact on self-care, occupational, and social activities. No specific measurement technology or method for determining the cutoffs is prescribed because this is a frame to capture multiple contexts of study. These 2 dimensions allow for 5 possible states represented in the oval shapes, including: acute low-impact pain and acute high-impact pain, chronic low-impact pain on chronic high-impact pain, and finally a special case of resolution of chronic pain. These 5 states allow for 10 possible transitions over time, which are given in Table 2. One cannot change from chronic to acute pain. This is a static representation of a set of dynamic processes, and we recognise that one can move between states over time, especially between chronic low impact and chronic high impact. We recognise also that chronic pain, once resolved, can relapse.

Table 2

Possible transitions between chronic pain states.

Chronic pain status	1st observation	2nd observation
Onset	Acute low impact	Chronic low impact
Onset	Acute low impact	Chronic high impact
Onset	Acute high impact	Chronic low impact
Onset	Acute high impact	Chronic high impact
Change (worsening)	Chronic low impact	Chronic high impact
Change (improving)	Chronic high impact	Chronic low impact
Change (resolution)	Chronic low impact	Chronic pain resolution
Change (resolution)	Chronic high impact	Chronic pain resolution
Maintenance	Chronic low impact	Chronic low impact
Maintenance	Chronic high impact	Chronic high impact

pain, a clinical diagnostic scheme that assumes high impact as a core part of the definition.³⁰

- (3) The concept of transition has been questioned because of an often tacit acceptance of a change in mechanism from acute to chronic pain. We make no reference to specific mechanisms but agree with the recommendation, where possible, "...to track individual pain types over time because they are likely to have different pain progression and resolution mechanisms and require different interventions."¹⁵
- (4) We also acknowledge that at any time point, acute and chronic pain can co-occur, but in this framework are focusing on the transition to/from chronic pain.
- (5) Related is the assumption that chronic pain is pain that is refractory to treatment. There is a need to establish evidence for refractory chronic pain monitoring treatment(s) and their unsuccessful outcomes. To date, there is no broadly accepted and generalisable definition of treatment-resistant chronic pain.
- (6) Similarly, Figure 1 is a static representation of changes in pain state. We recognise that one can move between high- and low-impact pain, and that pain can resolve and then relapse.⁴⁰
- (7) Living with longstanding pain can have a broad lasting impact on life that might endure past pain resolution.
- (8) A focus on duration and impact privileges the individual. Chronic pain has an impact beyond the individual to other individuals, to society and to the economy.

This duration-impact framework has been developed in the context of a major UK research programme investigating the psychosocial determinants of high-impact chronic pain⁷ funded by the Advanced Pain Discovery Platform.¹ The APDP has a focus on determining the causal influences on the onset and maintenance of high- and low-impact chronic pain. The consortium⁷ is exploring determinants of pain-state transitions using existing databases such as ALSPAC,^{4,18,32} ELSA,¹⁴ UK BIOBANK,⁴¹ HWW,²⁴ and HEAF²¹ the synthesis of findings across published studies,¹⁹ and through new investigations.

This framework is the first step in helping to clarify clinical and research questions. First, we need to understand how to manage the uncertainty inherent in the use of measurement technology designed to capture impact and establish how far what has already been measured corresponds with, or diverts from, this framework. Second, as we are interested in factors that are causally relevant to the onset, maintenance, and change in states over time, testing causal models needs to be carefully formulated. And third, this framework can direct the selection of appropriate endpoints for intervention in attempting to alter unwanted pain states. As important as pain offset (resolution) is the improvement in impact status, from high to low. A reasonable treatment outcome for many, and therefore a clinical endpoint, is to move from high-impact to low-impact chronic pain.^{10,11,36}

5. Conclusion

We propose a framework for studying the biopsychosocial influences on the onset, maintenance, and change in of chronic pain state. In accepting and interrogating the common dichotomies of duration (acute, chronic) and impact (high, low), we recognise the challenges inherent in dichotomizing continuous and dynamic experience. Pragmatically, however, this allows us to propose 5 unique states of pain and 10 transitions. This framework promotes a consideration of impact over time on the person with pain and will enable investigation of the causal determinants of states and changes in state.

Conflict of interest statement

The authors have no conflict of interest to declare.

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References

- Advanced pain discovery platform (APDP). Available at: https://www.ukri. org/what-we-offer/browse-our-areas-of-investment-and-support/ advanced-pain-discovery-platform-apdp/. Accessed January 27, 2023.
- [2] Boggini T, Pozzoli S, Schiavolin P, Erario R, Mosca F, Brambilla P, Fumagalli M. Cumulative procedural pain and brain development in very preterm infants: a systematic review of clinical and preclinical studies. Neurosci Biobehav Rev 2021;123:320–36.
- [3] Borsook D, Youssef AM, Simons L, Elman I, Eccleston C. When pain gets stuck: the evolution of pain chronification and treatment resistance. PAIN 2018;159:2421–36.
- [4] Boyd A, Golding J, Macleod J, Lawlor DA, Fraser A, Henderson J, Molloy L, Ness A, Ring S, Davey Smith G. Cohort profile: the 'children of the 90s'—the index offspring of the Avon longitudinal study of parents and children. Int J Epidemiol 2013;42:111–27.
- [5] Caeiro C, Moore A, Price L. Clinical encounters may not be responding to patients' search for meaning and control over non-specific chronic low back pain—an interpretative phenomenological analysis. Disabil Rehabil 2022;44:6593–607.
- [6] Cassel EJ. The nature of suffering and the goals of medicine. N Engl J Med 1982;306:639–45.
- [7] Consortium to research individual interpersonal and social factors in pain (CRIISP). Available at: https://www.bath.ac.uk/projects/consortium-toresearch-individual-interpersonal-and-social-influences-in-pain-criisp/. Accessed January 27, 2023.
- [8] Doshi TL, Dworkin RH, Polomano RC, Carr DB, Edwards RR, Finnerup NB, Freeman RL, Paice JA, Weisman SJ, Raja SN. AAAPT diagnostic criteria for acute neuropathic pain. Pain Med 2021;22:616–36.
- [9] Drissi I, Woods WA, Woods CG. Understanding the genetic basis of congenital insensitivity to pain. Br Med Bull 2020;133:65–78.
- [10] Dworkin RH, Turk DC, Farrar JT, Haythornthwaite JA, Jensen MP, Katz NP, Kerns RD, Stucki G, Allen RR, Bellamy N, Carr DB, Chandler J,

Cowan P, Dionne R, Galer BS, Hertz S, Jadad AR, Kramer LD, Manning DC, Martin S, McCormick CG, McDermott MP, McGrath P, Quessy S, Rappaport BA, Robbins W, Robinson JP, Rothman M, Royal MA, Simon L, Stauffer JW, Stein W, Tollett J, Wernicke J, Witter J; IMMPACT. Core outcome measures for chronic pain clinical trials: IMMPACT recommendations. PAIN 2005;113:9–19.

- [11] Dworkin RH, Turk DC, McDermott MP, Peirce-Sandner S, Burke LB, Cowan P, Farrar JT, Hertz S, Raja SN, Rappaport BA, Rauschkolb C, Sampaio C. Interpreting the clinical importance of group differences in chronic pain clinical trials: IMMPACT recommendations. PAIN 2009;146:238–44.
- [12] Eccleston C. Lives lived longer—chronic pain, subjective well-being, and occupation. In: Wainwright E, Eccleston C, editors. Work and pain: a lifespan developmental approach. Oxford: Oxford University Press, 2020. p. 91–102.
- [13] Eccleston C, Jordan AL, Crombez G. The impact of chronic pain on adolescents: a review of previously used measures. J Pediatr Psychol 2006;31:684–97.
- [14] English Longitudinal Study of Ageing (ELSA). Available at: https://www. elsa-project.ac.uk/. Accessed January 27, 2023.
- [15] Finnerup NB, Nikolajsen L, Rice ASC. Transition from acute to chronic pain: a misleading concept? PAIN 2022;163:e985–8.
- [16] Fordyce WE. Pain and suffering. A reappraisal. Am Psychol 1988;43: 276–83.
- [17] Fordyce WE. Pain and suffering: what is the unit? Qual Life Res 1994; 3(suppl 1):S51–6.
- [18] Fraser A, Macdonald-Wallis C, Tilling K, Boyd A, Golding J, Davey Smith G, Henderson J, Macleod J, Molloy L, Ness A, Ring S, Nelson SM, Lawlor DA. Cohort profile: the Avon longitudinal study of parents and children: ALSPAC mothers cohort. Int J Epidemiol 2013;42:97–110.
- [19] Gooberman-Hill R, Wainwright E, Guest A, Zeyen A, Sallis H, Stone S. Social and cultural influences in the experience of transition to or from long-term (chronic) pain: a meta-synthesis of qualitative research studies, PROSPERO 2022 CRD42022337979, 2022. Available from: https:// www.crd.york.ac.uk/prospero/display_record.php?ID=CRD42022337979
- [20] Headache Classification Committee of the International Headache Society (IHS). The international classification of headache disorders, 3rd edition. Cephalalgia 2018;38:1–211.
- [21] Health and Employment After Fifty (HEAF) Study. Available at: https:// www.mrc.soton.ac.uk/heaf/. Accessed January 27, 2023.
- [22] Herman PM, Qureshi N, Arick SD, Edelen MO, Hays RD, Rodriguez A, Weir RL, Coulter ID. Definitions of chronic low back pain from a scoping review, and analyses of narratives and self-reported health of adults with low back pain. J Pain 2023;24:403–12.
- [23] Hofer DM, Lehmann T, Zaslansky R, Harnik M, Meissner W, Stuber F, Stamer UM. Rethinking the definition of chronic postsurgical pain: composites of patient-reported pain-related outcomes vs pain intensities alone. PAIN 2022;163:2457–65.
- [24] Hurt L, Ashfield-Watt P, Townson J, Heslop L, Copeland L, Atkinson MD, Horton J, Paranjothy S. Cohort profile: HealthWise Wales. A research register and population health data platform with linkage to National Health Service Data Sets in Wales. BMJ Open 2019;9:e031705.
- [25] Ilhan E, Pacey V, Brown L, Spence K, van Ganzewinkel CJ, Pillai Riddell R, Campbell-Yeo M, Stevens BJ, Eriksson M, Shah V, Anand KJS, Bellieni C, Daly M, Johnston C, Hush J. What is the definition of acute episodic and chronic pain in critically ill neonates and infants? A global, four-stage consensus and validation study. BMJ Open 2022;12:e055255.
- [26] Interagency Pain Research Coordinating Committee. National Pain Strategy Report. Available at: https://www.iprcc.nih.gov/node/5/ national-pain-strategy-report. Accessed January 27, 2023.
- [27] Johansen A, Romundstad L, Nielsen CS, Schirmer H, Stubhaug A. Persistent postsurgical pain in a general population: prevalence and predictors in the Tromso study. PAIN 2012;153:1390–6.
- [28] Klint A, Bondesson E, Rasmussen BH, Furst CJ, Schelin MEC. Dying with unrelieved pain-prescription of opioids is not enough. J Pain Symptom Manage 2019;58:784–91.e1.
- [29] Kongsted A, Kent P, Axen I, Downie AS, Dunn KM. What have we learned from ten years of trajectory research in low back pain? BMC Musculoskelet Disord 2016;17:220.
- [30] Nicholas M, Vlaeyen JWS, Rief W, Barke A, Aziz Q, Benoliel R, Cohen M, Evers S, Giamberardino MA, Goebel A, Korwisi B, Perrot S, Svensson P, Wang SJ, Treede RD; IASP Taskforce for the Classification of Chronic Pain. The IASP classification of chronic pain for ICD-11: chronic primary pain. PAIN 2019;160:28–37.
- [31] Noel M, Chambers CT, Parker JA, Aubrey K, Tutelman PR, Morrongiello B, Moore C, McGrath PJ, Yanchar NL, Von Baeyer CL. Boo-boos as the building blocks of pain expression: an observational examination of

parental responses to everyday pain in toddlers. Can J Pain 2018;2: 74–86.

- [32] Northstone K, Lewcock M, Groom A, Boyd A, Macleod J, Timpson N, Wells N. The Avon Longitudinal Study of Parents and Children (ALSPAC): an update on the enrolled sample of index children in 2019. Wellcome Open Res 2019;4:51.
- [33] Pitcher MH, Von Korff M, Bushnell MC, Porter L. Prevalence and profile of high-impact chronic pain in the United States. J Pain 2019;20: 146–60.
- [34] Raja SN, Carr DB, Cohen M, Finnerup NB, Flor H, Gibson S, Keefe FJ, Mogil JS, Ringkamp M, Sluka KA, Song X-J, Stevens B, Sullivan MD, Tutelman PR, Ushida T, Vader K. The revised International Association for the Study of Pain definition of pain: concepts, challenges, and compromises. PAIN 2020;161:1976–82.
- [35] Ravn SL, Hartvigsen J, Hansen M, Sterling M, Andersen TE. Do posttraumatic pain and post-traumatic stress symptomatology mutually maintain each other? A systematic review of cross-lagged studies. PAIN 2018;159:2159–69.
- [36] Robinson ME, Brown JL, George SZ, Edwards PS, Atchison JW, Hirsh AT, Waxenberg LB, Wittmer V, Fillingim RB. Multidimensional success criteria and expectations for treatment of chronic pain: the patient perspective. Pain Med 2005;6:336–45.
- [37] Rome IV Diagnostic Criteria for Functional Gastrointestinal Disorders. Available at: https://theromefoundation.org/rome-iv/rome-iv-criteria/. Accessed January 27, 2023.
- [38] Treede RD, Rief W, Barke A, Aziz Q, Bennett MI, Benoliel R, Cohen M, Evers S, Finnerup NB, First MB, Giamberardino MA, Kaasa S, Korwisi B, Kosek E, Lavand'homme P, Nicholas M, Perrot S, Scholz J, Schug S, Smith BH, Svensson P, Vlaeyen JWS, Wang SJ. Chronic pain as a symptom or a disease: the IASP classification of chronic pain for the International Classification of Diseases (ICD-11). PAIN 2019;160: 19–27.
- [39] Turk DC, Fillingim RB, Ohrbach R, Patel KV. Assessment of psychosocial and functional impact of chronic pain. J Pain 2016;17:T21–49.
- [40] Turk DC, Rudy TE. Neglected topics in the treatment of chronic pain patients-relapse, noncompliance, and adherence enhancement. PAIN 1991;44:5–28.
- [41] UK Biobank. Available at: http://www.ukbiobank.ac.uk. Accessed January 27, 2023.
- [42] van den Berg-Emons RJ, Schasfoort FC, de Vos LA, Bussmann JB, Stam HJ. Impact of chronic pain on everyday physical activity. Eur J Pain 2007; 11:587–93.
- [43] Vlaeyen JWS, Haslbeck JMB, Sjouwerman R, Peters ML. Towards a dynamic account of chronic pain. PAIN 2022;163:e1038–9.
- [44] Vlaeyen JWS, Morley S, Crombez G. The experimental analysis of the interruptive, interfering, and identity-distorting effects of chronic pain. Behav Res Ther 2016;86:23–34.
- [45] Von Korff M, DeBar LL, Krebs EE, Kerns RD, Deyo RA, Keefe FJ. Graded chronic pain scale revised: mild, bothersome, and high-impact chronic pain. PAIN 2020;161:651–61.
- [46] Von Korff M, Dunn KM. Chronic pain reconsidered. PAIN 2008;138: 267–76.
- [47] Von Korff M, Scher AI, Helmick C, Carter-Pokras O, Dodick DW, Goulet J, Hamill-Ruth R, LeResche L, Porter L, Tait R, Terman G, Veasley C, Mackey S. United States national pain strategy for population research: concepts, definitions, and pilot data. J Pain 2016;17:1068–80.
- [48] Vowles KE, Rosser B, Januszewicz P, Morlion B, Evers S, Eccleston C. Everyday pain, analgesic beliefs and analgesic behaviours in Europe and Russia: an epidemiological survey and analysis. Eur J Hosp Pharm 2014; 21:39–44.
- [49] Wall PD. On the relation of injury to pain. The John J. Bonica Lecture. PAIN 1979;6:253–64.
- [50] Walters ET, Williams ACdC. Evolution of mechanisms and behaviour important for pain. Philos Trans R Soc B Biol Sci 2019;374:20190275.
- [51] Weisman A, Quintner J, Masharawi Y. Congenital insensitivity to pain: a misnomer. J Pain 2019;20:1011–4.
- [52] Werner MU, Kongsgaard UEI. Defining persistent post-surgical pain: is an update required? Br J Anaesth 2014;113:1–4.
- [53] Williams ACdC. Persistence of pain in humans and other mammals. Philos Trans R Soc B Biol Sci 2019;374:20190276.
- [54] Williams ACdC, Fisher E, Hearn L, Eccleston C. Psychological therapies for the management of chronic pain (excluding headache) in adults. Cochrane Database Syst Rev 2020;8:CD007407.
- [55] Yong RJ, Mullins PM, Bhattacharyya N. Prevalence of chronic pain among adults in the United States. PAIN 2022;163:e328–32.

5