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Citation for final published version:

Smith, William Daniel, Maier, Wolfgang D. and Bliss, Ian 2020. Contact-Style magmatic sulphide mineralisation in the Labrador Trough, northern Québec, Canada: Implications for regional prospectivity. Canadian Journal of Earth Sciences 57 (7), pp. 867-883. 10.1139/cjes-2019-0137

Publishers page: http://dx.doi.org/10.1139/cjes-2019-0137

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- 1 Contact-style magmatic sulphide mineralisation in the Labrador Trough, northern
- 2 Québec, Canada: Implications for regional prospectivity
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## Abstract

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- 8 The Labrador Trough in northern Québec is currently the focus of ongoing exploration for
- 9 magmatic Ni-Cu-PGE sulphide ores. This geological belt hosts voluminous basaltic sills and
- 10 lavas of the Montagnais Sill Complex, which are locally emplaced amongst sulphidic
- metasedimentary country rocks. The recently discovered Idefix PGE-Cu prospect represents a
- stack of gabbroic sills that host stratiform patchy net-textured sulphides (0.2 to 0.4 g/t PGE +
- Au) over a thickness of  $\sim 20$  m, for up to 7 km. In addition, globular sulphides occur at the
- base of the sill, adjacent to the metasedimentary floor rocks. Whole-rock and PGE
- 15 geochemistry indicates that the sills share a common source and that the extracted magma
- underwent significant fractionation before emplacement in the upper crust. To develop the
- 17 PGE-enriched ores, sulphide melt saturation was attained before final emplacement, peaking
- at R factors of  $\sim 10,000$ . Globular sulphides entrained along the base of the sill ingested
- crustally-derived arsenic and were ultimately preserved in the advancing chilled margin.
- 20 **Keywords:** Labrador Trough, PGE, magmatic sulphide, mineral exploration

#### 1. Introduction

The Montagnais Sill Complex (MSC) in the Labrador Trough of northern Québec is currently the focus of ongoing exploration for magmatic Ni-Cu-platinum group element (PGE) sulphide deposits. The MSC is considered prospective because it hosts voluminous basaltic rocks (> 10,000 km²), in proximity to a rifted craton margin (*e.g.*, Begg *et al.* 2010), locally emplaced amongst sulphidic metasedimentary rocks (*e.g.*, Ripley and Li 2013). Despite the apparent prospectivity of the province, exploration remains in its infancy, hindered by a lack in understanding of the ore-forming processes operating in the region. Recent exploration conducted by Northern Shield Resources has identified numerous prospective showings across the Labrador Trough, including the Idefix PGE-Cu prospect.

The Idefix PGE-Cu prospect is located approximately 75 km west of Kuujjuaq (Fig. 1) and encompasses 41 claims over 18.5 km². The prospect comprises a sequence of differentiated gabbroic sills, bound by intercalated metapelites of the Baby Formation. To date, 1501 m of diamond drilling across fourteen boreholes has revealed stratiform patchy net-textured sulphide ore, with an average 2PGE (Pt + Pd) grade of 0.2 to 0.4 g/t over a thickness of 16 to 34 m, traceable for approximately 7 km along strike. In this paper, we examine the architecture of the Idefix prospect and investigate the processes that led to its formation.

## 2. Regional Setting

The Labrador Trough represents a foreland fold-and-thrust belt, extending for ~ 800 km from the Grenville Front to Ungava Bay (Hoffman 1988; Skulski *et al.* 1993). The trough formed during the oblique collision between the Southern Rae Craton and the Superior Craton at 1.84 to 1.82 Ga (Wares and Goutier 1990; Wardle and Van Kranendonk 1996). This collision

instigated pervasive, westward out-of-sequence imbricate thrusting of thick-skinned passive margin sediments, where thrust zones are interpreted to propagate to the base of the crust (Wares and Goutier 1990). Metamorphism increases from west to east, peaking at greenschist facies in the northwest (Perreault and Hynes 1990)

Previous work characterised the stratigraphy of the Labrador Trough using a cyclic system, where cycles 1 and 2 represent episodes of passive margin sedimentation, topped by a third cycle of flysch deposits (Clark and Wares 2005 and references therein). Basalts and subordinate rhyolites were emplaced during cycle 1 at  $2169 \pm 4$  Ma and  $2142 \pm 4$  Ma, respectively, marking the onset of rift-related magmatism (Skulski *et al.* 1993 and references therein). The MSC intruded into sedimentary rocks of cycles 1 and 2 at  $1884 \pm 1.6$  Ma (Findley *et al.* 1995). Co-magmatic basalts of the Willbob and Hellancourt Formations (in the south and north, respectively) were emplaced contemporaneously with the MSC (Rohon *et al.* 1993). Several hypotheses for the tectonomagmatic setting of the MSC have been proposed, including: (1) dextral oblique extension above an east-dipping subduction zone (Hoffman 1990b), (2) dextral transtension along the eastern margin of the Superior Craton, resulting in the formation of pull-apart rift basins (Skulski *et al.* 1993), (3) extension in an ensialic backarc setting (Corrigan *et al.* 2009), and (4) derivation from a deep-seated mantle plume (Ciborowski *et al.* 2017).

The MSC comprises three major sill types, including (i) aphyric (equigranular) mesogabbros with stratiform gabbroic pegmatites, (ii) glomeroporphyritic and porphyritic gabbros, and (iii) differentiated gabbro sills, sometimes with a basal peridotite cumulate layer. Each of these sill types has been described as hosting magmatic sulphide mineralisation (Clark and Wares 2005). Mineralised sills are concentrated in the upper part of the cycle 2 sedimentary package, which comprises sulphide-bearing sediments of the Baby (in the north)

and Menihek Formations (in the south), suggesting a link between crustal contamination and sulphide melt saturation in the magmas (Clark and Wares 2005).

Grab and channel samples were collected across the Idefix property during field

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#### 3. Methods

reconnaissance in 2012 and 2013. Additional drill core was acquired from the headquarters of Northern Shield Resources, generated during their 2013 drilling program. Thick and thin sections were petrographically analysed and photomicrographed using a Leica MZ12s microscope with camera attachment at Cardiff University. In total, sixty whole-rock samples were analysed for lithophile major and trace elements, whereas 706 whole-rock samples were assayed for major lithophile elements, V, Cr, Co, Ni, Cu, Zn, Sr, Pt, Pd, and Au. Twenty-eight whole-rock samples from Idefix property and the surrounding gabbroic rocks were analysed for a full suite of platinum group elements (PGE). All geochemical analyses were performed by ALS Minerals (Vancouver). Sample preparation was completed at ALS Minerals (Timmins) using the PREP-31 package. Lithophile major and trace elements were determined through ICP-AES and ICP-MS respectively, following four-acid digestion of fused beads (ALS codes ME-ICP06, ME-MS81, ME-MS42\*, ME-4ACD81, and ME-ICP61a). Loss on ignition (LOI) was determined using the OA-GRA05 package. Sulphur and carbon concentrations were determined using a Leco sulphur analyser, whereby 0.1 g of homogenised sample is combusted at ~ 1350°C and total S and C are measured using a non-dispersive infrared sensor (ALS code ME-IR08). Palladium, Pt, and Au concentrations were determined by ICP-MS, following lead oxide fire assay to produce a precious metal bead (ALS codes FA-FUSPG4 and PGM-MS23). To

determine the full suite of six PGE and Au, a precious metal bead was produced via nickel

sulphide fire assay and measured through ICP-MS (ALS codes FA-FUSNS01 and PGM-MS25NS). More information regarding analytical procedures can be found on the ALS Minerals website (<a href="www.alsgobal.com">www.alsgobal.com</a>). Results for representative samples are presented in Tables 1 and 2. The full dataset, as well as standards and blanks, is reported in Supplementary Material 1.

A Zeiss Sigma HB Field Emission Gun Analytical Scanning Electron Microscope, equipped with two Oxford Instruments 150 mm<sup>2</sup> energy dispersive spectrometers (EDS) was used to generate element maps of the sulphide-bearing rocks. The maps were produced at a working distance of 8.9 mm, an accelerating voltage of 20 kV, and a pixel dwell time of 2000 ms.

## 4. The Idefix PGE-Cu Prospect

## 4.1. Historical exploration

Airborne reconnaissance surveys conducted in the 1940s identified numerous gossanous outcrops in the broader area hosting Idefix, which were later identified to be magmatic sulphide showings. In the 1950s, several companies conducted field reconnaissance and geophysical surveys in the vicinity of the Idefix prospect and identified several Ni-Cu showings (Sogemines Development Company 1962; Schilling 1963; De Freneuse 1965). Cities Service Minerals Corporation reported the first PGE values approximately ~ 5 km west of Idefix, in gabbroic rocks grading 10.3 g/t Pd, 3.4 g/t Pt, 11.2% Cu, and < 0.1% Ni (Loring 1975).

La Fosse Platinum Group conducted extensive Ni-Cu-PGE exploration along the eastern half of the Labrador Trough during the 1980s and were the first to identify mineralisation at what is now Idefix (then named Ukunngaalik). Samples of aphyric

mesogabbro returned grades of 5.8 g/t Pd, 2.7 g/t Pt, and 0.8 g/t Au (Avison 1986; La Fosse Platinum Group 1988). The area was not explored further until 2002 when Virginia Mines Incorporated identified additional stratiform sulphide mineralisation with grades of up to 18.1 g/t 2PGE + Au (Savard 2003).

Northern Shield Resources conducted seasonal field surveys across the eastern half of the Labrador Trough from 2011 to 2013. Out of 1,614 samples collected from the region, 934 assayed > 0.1 g/t PGE. The Idefix prospect was staked in 2011 and encompasses 41 claims covering 18.5 km<sup>2</sup>. Fourteen diamond-drill holes totalling 1,501 m were drilled along the Idefix ridge (Fig. 1), in which grades of 0.2 to 0.4 g/t 2PGE (Pt and Pd) over 16 to 34 m can be traced intermittently for approximately 7 km.

## 4.2. Local Geology

The Idefix prospect is located within the Gerido lithotectonic zone defined by Clark and Wares (2005; Fig. 1b). The Gerido zone is characterised by intercalated metasediments, banded iron formations, and volcanic extrusives, intruded by mesogabbroic and ultramafic sills of the MSC. Fine- to medium-grained equigranular mesogabbroic rocks strike NW-SE, with dips varying from 85° to 40° to the east (Fig. 1c).

Borehole 13ID-13 (Fig. 2a) intersects the entire stratigraphy of the Idefix property and has been used to characterise the stratigraphy of the prospect. The country rock consists of stratified metapelites, locally containing subhedral pyrite (Fig. 2b). The contact between the country rock and the composition Idefix sill is sharp and irregular, sometimes characterised by a < 5 cm thermal aureole. The Idefix sill is  $\sim$  100 m thick and has been sub-divided into I1, I2, and I3, based on the presence of sulphide mineralisation, as outlined in detail below.

The Idefix sill is overlain by a further ~ 400-m-thick mesogabbroic hanging wall sill named the Primitive Unit (PU). The PU contains only sparsely disseminated sulphide mineralisation.

Relatively thin stratiform gabbroic pegmatite horizons (< 0.5 m) intermittently occur along the I1-I2 and I3-PU contacts (Fig. 2c-d). The former is typically less defined than the latter, but where present, both contain finely disseminated sulphide mineralisation and bluish, interstitial quartz.

# 4.3. Petrology of the Idefix and Primitive aphyric gabbro sills

Gabbroic rocks of the Idefix prospect are fine- to medium-grained and composed of partially uralitized anhedral clinopyroxene and partially to completely saussurised subhedral plagioclase (Fig. 3). There is no evidence for serpentinization of olivine in any samples and no discernible deformation. The PU is characterised by a higher relative proportion of clinopyroxene (~ 53 vol.%) at the expense of plagioclase (~ 45 vol.%; Fig. 3a). The Idefix sills generally comprise a higher proportion of modal plagioclase (~ 48 vol.%) than clinopyroxene (~ 46 vol.%; Fig. 3b). Accessory phases (< 2 vol.%) include hornblende, acicular actinolite and tremolite, biotite, anhedral apatite, titanite, and Fe-Ti oxides. Anhedral apatite and clusters of titanite, biotite, Fe-Ti oxide, and rutile are most prevalent in sulphidebearing samples. The I1 sill differs from the other Idefix sills in that it can locally host up to ~ 3 vol.% interstitial quartz. CIPW normative mineralogy of the gabbroic pegmatites indicates that there is little difference relative to that of the aphyric gabbroic sills. The I3-PU gabbroic pegmatite comprises up to ~ 8 vol.% interstitial quartz, with traces of apatite, biotite, ilmenite, and titanite. The I1-I2 pegmatite is characterised by up to ~ 17 vol.% interstitial quartz (Fig. 3c).

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Two horizons of mineralisation have been intersected at Idefix: (i) patchy net-textured sulphides (< 5 vol.%; Fig 4a) in the contact zone between I3 and PU, and (ii) globular sulphides (0.5 to 2 cm in diameter) in the basal chilled margin of the Idefix sill (I1; Fig. 4b). Finely disseminated sulphide (< 1 vol.%) is observed throughout the remainder of the Idefix sill (*e.g.*, Giles 2015).

The patchy net-textured ores in the I3-PU contact zone are dominated by pyrrhotite  $[Fe_{(1-x)}S; Po; \sim 60 \text{ vol.\%}]$ , chalcopyrite (CuFeS<sub>2</sub>; Ccp;  $\sim 32 \text{ vol.\%}$ ), with subordinate pentlandite  $[(Fe,Ni)_9S_8; Pn; \sim 8 \text{ vol.\%}]$ . Sparsely disseminated sulphide throughout the remainder of I2 and I3 is generally dominated by Ccp over Po, with little-to-no Pn. Base metal sulphides are spatially associated with apatite ( $\sim 0.2 \text{ vol.\%}$ ) and titanite ( $\sim 0.3 \text{ vol.\%}$ ).

Sulphide globules in I1 are often elliptical and comprise a Po core (~ 62 vol.%), enclosing granular and flame Pn (~ 3 vol.%). Chalcopyrite (~ 33 vol.%) occurs around the exterior of the globules and is disseminated throughout the unit. Chalcopyrite is often spatially associated with sulpharsenides belonging to the cobaltite-gersdorffite series (~ 1 vol.%), sphalerite (< 0.1 vol.%), titanite (< 1 vol.%) and apatite.

# 5. Results

## 5.1. Lithophile elements

All gabbroic rocks at Idefix plot within a narrow range of MgO (~ 8 to 14 wt.%; Table 1), in that the PU is generally more primitive (10.4 to 13.9 wt.%) than the Idefix gabbros (8.4 to 12.7 wt.%). Throughout all samples, CaO and Cr/V decrease with decreasing MgO content and FeOt, TiO<sub>2</sub> and Sr increase with decreasing MgO content (Fig. 5a-d). The incompatible element composition of the PU and I3 are generally similar (*e.g.*, TiO<sub>2</sub>, K<sub>2</sub>O, Sr, and REE),

whereas I2 and I1 show greater enrichment in these elements (Fig. 5e). The greatest spread of data is observed in Cr/V values (Fig. 5d), notably for the PU (Cr/V ~ 2 to 6.5). The average Cr/V content of the Idefix sub-units decreases from I3 (Cr/V 1.9 to 6.3) to I1 (Cr/V 0.8 to 2.9). The majority of samples show a Ni concentration of 200 to 250 ppm. A few samples, notably from I3, show higher Ni (> 300 ppm), despite no variation in MgO content. The high Ni contents correspond to an increase in sulphide content.

Primitive mantle normalised multi-element plots (Fig. 6) show broadly similar profiles for gabbroic rocks. The patterns of the PU and I3 are relatively flat (1 to 2x primitive mantle), with variably negative P anomalies. Sub-units I1 and I2 show slightly enriched patterns (2 to 4x primitive mantle), with less pronounced negative P anomalies and strong positive K anomalies. Local metapelitic rocks of the Baby Formation show strong enrichment in LILEs and LREEs, but a similar HREE level relative to that of the gabbroic units. All gabbroic units are characterised by similar Th/Yb<sub>N</sub> (0.4 to 1.1), La/Sm<sub>N</sub> (0.8 to 1.2), and Gd/Yb<sub>N</sub> (0.8 to 1.2) ratios, whereas metasedimentary units have Th/Yb<sub>N</sub> values > 50 and La/Sm<sub>N</sub> values > 4.8. Total REE concentration increases with decreasing MgO. All gabbroic units show sub-parallel REE patterns, with weakly positive or negative Eu anomalies (Eu/Eu\*  $\sim$  0.8 to 1.2; Table 1).

# 5.2. *Chalcophile elements*

In all gabbroic samples, chalcophile metal concentrations increase with increasing sulphur (Table 2; Fig. 7a-c). The patchy net-textured ores (I3) are generally more enriched in PGE relative to globular ores (I1; Fig. 7c). This trend may correspond to a difference in the platinum group mineral assemblage of the two ore-bearing horizons. Platinum and Pd show a good positive correlation across both ore-bearing horizons ( $R^2 = 0.89$ ), with an average Pd/Pt

ratio of  $\sim 2.8$  (Fig. 7d). The base metals Cu and Ni show good positive correlations with PGE, with the patchy net-textured ores being relatively more enriched in PGE (Fig. 7e-f).

Nickel, the IPGE (Ir and Ru), and Rh show good positive correlations across all ore-bearing gabbroic rocks (Fig. 8a-c). Regional gabbro (RG) samples represent mineralised mesogabbroic rocks at or near to the Idefix property. All samples have Pd/Ir values between 200 and 500 (Fig. 8d), within the range of basaltic rocks elsewhere (e.g., Maier et al. 2008b). Platinum and Au contents broadly increase with increasing Ir (Fig. 8e-f). All ore-bearing rocks at Idefix possess Ni/Ir<sub>N</sub> and Cu/Pd<sub>N</sub> values < 1 (Fig. 9). All samples (> 0.5 wt.% S) plotted in Figure 9 have been recalculated to 100% sulphide using the method of Barnes and Lightfoot (2005). Both ore types exhibit similar patterns, in that IPGE levels are ~ 100x that of the primitive mantle, whereas PPGE (Rh, Pt, and Pd) levels are 1,000 to 10,000x that of the primitive mantle. Globular ores are more enriched in Au and Cu, relative to patchy nettextured ores. The Idefix profiles are similar to that of the Paladin deposit in the Labrador Trough (see Clark and Wares 2005), which represents PGE-enriched sulphide mineralisation in an aphyric gabbro with stratiform gabbroic pegmatite (deposit type 10d in Clark and Wares, 2005). They also show some similarity to the pattern of the J-M Reef, in that the rocks ores are low in IPGE, high in PPGE, with negative Au anomalies.

Figure 10 shows that the majority of sulphide-bearing gabbroic rocks have Cu/Pd ratios below the range of the primitive mantle. This suggests that the host magma did not lose significant sulphide melt before its final emplacement, which would have resulted in a marked depletion in PGE relative to base metals. Overlain on this plot are *R* factor estimations assuming a parent magma composition similar to high-MgO basalts from the Cape Smith Belt (Barnes *et al.* 1992). Most samples plot between an estimated *R* factor range of 1,000 to 10,000 at < 1 vol.% sulphide, consistent with sulphide volumes observed in rock samples.

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#### 5.3. Borehole 13ID-13

The borehole samples the hanging wall PU (15 to 115 m), the Idefix sill (115 to 205 m), and metasedimentary footwall lithologies (205 to 206 m). The latter are marked by a pronounced decrease in MgO, Mg#, and Ni (Fig. 11). The composition of I1 becomes less evolved with height, as reflected by an upward increase in MgO, Mg#, and Cr/V, and a decrease in TiO<sub>2</sub> and Sr. In the contact zone between I1 and I2 is a quartz-bearing intermittent gabbroic pegmatitic horizon, which shows little change in composition with the bounding sub-units. Upward through I2, MgO and Cr/V gradually decrease, whereas S and chalcophile metals (Ni < Cu) increase. From the contact between I2 and I3 upwards, MgO and Cr/V increase toward the PU, whilst TiO<sub>2</sub> and Sr gradually decrease. Similarly, S decreases with height, while chalcophile metals increase, peaking at the contact between I3 and the PU. Mafic and incompatible elements smoothly increase and decrease respectively, into the PU. Chalcophile metal concentrations in the PU are mostly at or close to the detection limit. There is little-tono variation in Cu/Pd and Pd/Pt values upward through I1 and I2. I3 is characterised by lower Cu/Pd values (< 2,000) and upward decreasing Pd/Pt values. The PU comprises generally higher Cu/Pd values and upward decreasing Pd/Pt values (Fig. 11). These vertical patterns are observed in other drilled boreholes in the property (Supplementary Figure 1).

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## 6. Discussion

# 6.1. Interpreting the stratigraphic orientation

The orientation of the mafic rocks is of major importance in designing effective ore targeting models. However, trends in whole-rock geochemistry at Idefix are inconclusive. Most differentiation indices (Mg#, Cr/V, Ti, Pd/Ir, and Cu/Pd) show a broad trend towards

relatively unevolved compositions with height across the Idefix sill. While this could suggest that the Idefix sill has been tectonically overturned, it is also known that many mafic intrusions show basal reversals on the scale of meters to hundreds of meters in relation to a progressively decreasing trapped liquid component with height in the sill (*e.g.*, Wingellina Hills; Maier *et al.* 2015).

The Idefix PGE-Cu property belongs to deposit type 10d described by Clark and Wares (2005). Several similarities exist between the Idefix prospect and previously discovered occurrences, including the Paladin, Lac Lafortune, and Lac Larochelle prospects. All of these occurrences reside in the Gerido lithotectonic zone in the northern half of the Labrador Trough and are enriched in PGE relative to base metals. Furthermore, sulphide mineralisation (< 5 vol.%) at these occurrences is confined to stratiform gabbroic pegmatites and the underlying gabbro (Clark and Wares 2005 and references therein). Clark and Wares (2005) state that basal sulphides are not observed in these deposit types and that the stratiform gabbroic pegmatitic horizons represent the roof of the sill. However, globular sulphide is present at the base of the Idefix gabbro (with seemingly no compromise to the Cu/Pd values of the overlying net-textured sulphide) and there is evidence for two stratiform gabbroic pegmatites (*i.e.*, 11-12 and I3-PU). This could either mean that stratiform gabbroic pegmatites may not be exclusive to the upper portions of the sills or that the Idefix sub-units are all distinct sub-sills.

Most PGE-Cu reefs (*e.g.*, Bushveld Complex, Maier *et al.* 2008 and Penikat, Iljina *et al.* 2015) display a sharp basal peak in PGE concentrations, which is overlain by exponentially decreasing PGE over distances varying from a few decimetres to metres. This is typically interpreted to reflect Rayleigh-type fractionation of PGE in an S-saturated magma (Naldrett *et al.* 2009). However, in few PGE-Cu reefs, PGE contents decrease downward into the footwall rocks (*e.g.*, Merensky Reef, Barnes and Maier 2002; Stillwater Complex, Godel

2007). In the Merensky Reef, this has been ascribed to downward percolation of sulphide liquid into semi-consolidated floor rocks (Barnes and Maier 2002). No obvious systematic change in Cu/Pd and Ni/Cu values preclude the percolation of progressively fractionating sulphide liquid and instead suggests that gravity-driven percolation is responsible for the downward decreasing tails of chalcophile metals observed at Idefix (Mungall and Su 2005).

Sulphide liquid cannot travel for significant distances without dispersing into sub-millimetre disseminations induced by Rayleigh-Taylor instabilities or by dissolving in a depressurising magma (Mavrogenes and O'Neill 1999; Robertson *et al.* 2015). Due to the high density of sulphide melt, it tends to be concentrated along the base of magma bodies. However, globular sulphide documented at Noril'sk-Talnakh are an exception (Arndt *et al.* 2003). These globules are thought to have been transported upward in the host sill by gas bubbles, evidenced by the presence of silicate caps atop sulphide globules (see Barnes *et al.* 2019). Sulphide globules at Idefix do not possess silica caps, nor do they possess a Po-Pn lower margin and a Ccp upper margin, which has been used as geopetal structures for host rocks elsewhere (Prichard *et al.* 2004). However, granular pentlandite between chalcopyrite and pyrrhotite supports *in situ* fractionation for the sulphide globules (*e.g.*, Mansur *et al.* 2019). The occurrence of globules in I1 favours an up-right stratigraphy if this sub-unit represents the base of the Idefix or Primitive sills.

## 6.2. Sulphide melt saturation

Primitive magmas are typically undersaturated in sulphide melt as they ascend through the crust, due to the inverse relationship between sulphide melt solubility and pressure (*e.g.*, Mavrogenes and O'Neill 1999). As the magma undergoes differentiation, the sulphur content of the melt increases as a result of its incompatibility within silicate minerals. For basaltic

magmas, sulphide melt saturation can be attained either through extensive low-pressure fractional crystallisation (typically 40 to 60%) and/or through assimilation of exogenous sulphur (Mavrogenes and O'Neill 1999; Ripley and Li 2013). In many world-class Ni-Cu-PGE sulphide deposits, the assimilation of sulphur-rich country rock is considered a key process in triggering sulphide segregation (Kabanga, Maier *et al.* 2010; Noril'sk-Talnakh, Ripley *et al.* 2003; Voisey's Bay, Li and Naldrett 1999). However, this process is not necessarily required in the formation of PGE-Cu deposits. Instead, it is argued that in these deposit types, fractional crystallisation drives the magma to sulphide melt saturation, resulting in low volumes of sulphide (~ 1%) and higher *R* factors (> 10,000; Campbell and Naldrett 1979). A third possibility is that the Idefix magmas assimilated and/or entrained proto-ore from antecedent pulses of magma (*i.e.*, PU; Maier 2005; Maier and Groves 2011).

Firstly, there is no evidence that the host magmas at Idefix underwent crustal assimilation (Fig. 12). Binary mixing diagrams between a parent magma with the composition of the co-magmatic Hellancourt basalt (Ciborowski *et al.* 2017) and the local Baby Formation metasediments show that all gabbroic rocks (including the ore-bearing horizons) have assimilated < 5% of the country rock (Fig. 12a-b). This conclusion is consistent with those made for sulphide occurrences in the south of the Labrador Trough (*e.g.*, Chauval *et al.* 1987; Rohon *et al.* 1993). It is also consistent with S/Se ratios that are widely used to constrain the presence of crustal sulphur in magmatic sulphides (Queffurus and Barnes 2015). This is because crustal sulphides typically possess much higher S/Se values than magmatic sulphide. At Idefix, all gabbroic rocks plot either at or below the estimated S/Se mantle range of 2850 to 4300 (Eckstrand and Hulbert 1987). However, country rocks are characterised by low S/Se values (< 2,000). Low S/Se values in the gabbroic rocks are consistent with an interaction between sulphide and S-undersaturated fluids (Queffurus and Barnes 2015). Little-to-no change in La/Sm<sub>N</sub> relative to S/Se values

argues against wholesale assimilation of country rock, but can be reconciled with devolatilisation of the pyrite-bearing country rock (*e.g.*, Ripley 1981). Addition of exogenous sulphur and arsenic via country rock devolatilisation to I1 may explain the lower PGE tenors and increased relative proportion of sulpharsenides of the globular ores (*e.g.*, Piña *et al.* 2013; Le Vaillant *et al.* 2018), yet this cannot explain low S/Se values in I3 due to the proximity of this unit to the floor rocks.

For the estimation of the R factor, we assume that the parent magma to the Idefix gabbro underwent closed-system fractional crystallisation (Campbell and Naldrett 1979). The Cu concentration is within the Cu range of the co-magmatic Hellancourt basalts of Ciborowski *et al.* (2017;  $116 \pm 34$  ppm). From this, all gabbroic rocks at Idefix plot at approximate R factors of 5,000 to 10,000, at a maximum of  $\sim$  1 vol.% sulphide, which is consistent with sulphide proportions observed in drill-core and petrographically.

With no indication of crustal contamination influencing sulphide segregation, one must conclude that the Idefix magma was driven to sulphide melt saturation through differentiation. Clark and Wares (2005) argue that the relationship between disseminated sulphide and gabbroic pegmatites favours a hydrothermal origin. However, Pd/Ir ratios at Idefix are < 500 and there is no prominent negative Pt anomaly (e.g., Barnes and Liu 2012). The Cu/Pd values suggest that no PGE has been lost before the final emplacement of the Idefix magma, yet if sulphide segregated *in situ* after ~ 40 to 60% fractional crystallisation (*i.e.*, Ripley and Li 2013) there would be insufficient PGE-undepleted magma for immiscible sulphide to interact with. It must then be concluded that the Idefix magma became saturated in sulphide melt before its final emplacement (*i.e.*, magma conduit), perhaps aided by the cannibalisation of proto-ore left by antecedent pulses of magma (e.g., Maier, 2005; Maier and Groves 2011).

6.3. Assessing pre-emplacement fractionation of the Idefix magma

The enrichment in strongly chalcophile metals at Idefix suggests that the parent magma was undepleted in chalcophile metals and was derived from a large degree mantle melt. The flat REE profiles (La/Gd<sub>N</sub> = 0.8 to 1.2) of the gabbroic rocks at Idefix are analogous to REE profiles of co-magmatic Hellancourt basalts (Skulski *et al.* 1993; Rohon *et al.* 1993), which, in turn, resemble modern transitional MORB. Olivine (Fo ~ 0.84) is typically the liquidus phase in the Hellancourt basalts (Ciborowski *et al.* 2017). As olivine is not present in the Idefix gabbroic rocks, it is possible that the magmas underwent pre-emplacement fractional crystallisation. Pre-emplacement olivine fractionation would deplete the remaining melt of Ni. However, the gabbroic rocks at Idefix retain Ni values consistent with local basaltic rocks (~ 133 ppm). This is contradicted by the absence of positive Eu anomalies, which is to be expected if the magma had fractionation plagioclase prior to its final emplacement.

Moreover, the negative Ru anomalies may be best explained by pre-emplacement fractionation of spinel (Righter *et al.* 2004; Pagé and Barnes 2012). It is therefore, possible that the magma parental to the Idefix sills and associated basaltic rocks underwent pre-emplacement fractionation in the lower crust (*e.g.*, Skulski *et al.* 1993; Heinonen *et al.* 2019).

6.4. Emplacement and orientation of the gabbroic rocks at Idefix

From the current data, it remains unclear if the strata at Idefix has been tectonically overturned or not (see Fig. 13). The presence of two distinct sulphide-bearing horizons suggests at least two episodes of sill emplacement has occurred at Idefix, yet the sequence of emplacement is difficult to discern:

Scenario 1: Sequentially emplaced and overturned:

In this scenario, the PU was first emplaced, followed by 13 + 12, and then I1. To constrain the way-up of the PU, the eastern contact of this unit must be characterised. From the current data, there is no evidence that the PU overlying the Idefix sill is the upper part of a differentiating sill (MgO > 10 wt.%, TiO<sub>2</sub> < 0.5 wt.%). Although, it remains possible that this could represent a reversal if emplaced against the country rock. The Idefix sill was emplaced, whilst entraining immiscible sulphide, which underwent gravity-driven percolation to begin to accumulate on the I3-PU contact (Mungall and Su 2005). Lastly, I1 was emplaced entraining immiscible sulphide globules. However, if emplaced last, antecedent pulses of magma would have already devolatilised the country rock, precluding the formation of sulpharsenides by ingestion of crustally derived As. In this model, the gabbroic pegmatites form via static recrystallisation as described by Barnes and Maier (2002). The sequence was then tectonically overturned during the waning stages of the New Québec Orogeny. However, current mapping data support a regionally NE-facing sequence (M. Houlé, written communication, 2019),

## Scenario 2: Non-sequentially emplaced and overturned:

In this scenario, PU and I1 were emplaced before I2 + I3, whereby I1 represents the upper chilled margin of the PU. This is consistent with the generally chalcophile-depleted character of the PU ( $Cu/Pd \ge 10,000$ , PGE < 0.1 ppm). The density of sulphide melt typically precludes their occurrence in the upper parts of host sills, yet this has been documented elsewhere (*i.e.*, Noril'sk-Talnakh) due to volatile-rich bubbles (Barnes *et al.* 2019). However, the absence of silica caps around the globules, together with the high settling velocities of sulphide liquid (Chung and Mungall 2009) makes this unlikely at Idefix. The Idefix magmas were then emplaced below the advancing chilled margin, where entrained sulphide percolated to the

base of the sill. The I3-PU gabbroic pegmatite would have formed via static recrystallisation, yet the formation of the I1-I2 pegmatite may have been influenced by magmatic fluids exsolved from the Idefix magma. However, the current data cannot support this hypothesis.

# Scenario 3: Sequentially emplaced:

In this scenario, each sill represents an individual pulse of magma. Firstly, I1 was emplaced entrained immiscible sulphide globules that ingested crustally derived S and As. Secondly, I2 was emplaced, forming a discontinuous gabbroic pegmatite via static recrystallisation.

Thirdly, I3 was emplaced, whilst entraining PGE-rich immiscible sulphide. A gabbroic pegmatite has not been identified between I2 and I3, meaning that these units could represent one pulse of magma. Lastly, the PU was emplaced above the Idefix sill. The I3-PU gabbroic pegmatite is better preserved than the one present at the I1-I2 contact since the PU is thicker and less evolved than the Idefix sill.

## Scenario 4: *Non-sequentially emplaced*:

In this scenario, PU and II were emplaced before the Idefix sills, similar to that described in Scenario 2. This model can be reconciled with the (i) low PGE concentrations of the overlying PU and (ii) the assimilation of exogenous S and As from the country rock floor. The Idefix sill was progressively emplaced above the advancing chilled margin of the PU (*i.e.*, II). The I2 and I3 units may have been emplaced as two separate pulses, in that the I3 unit entrained a higher volume of immiscible sulphide as evidenced in the drill-core geochemistry (Fig. 11; Supplementary Figure 1).

# 6.5. Potential for PGE-Cu deposits in the Labrador Trough

In the Labrador Trough, PGE-Cu deposits are exclusive to aphyric gabbro sills in the Gerido lithotectonic zone (*e.g.*, Paladin, Lac Lafortune, and Lac Larochelle; Clark and Wares 2005). Their enrichment in PGE (Cu/Pd < 10,000) indicates that the parental magmas were undersaturated in sulphide melt during their emplacement. This suggests that they could have produced economic contact-style mineralisation. This deposit type is not necessarily dependent on the assimilation of crustal sulphur. Country rock in proximity to the Idefix sill is only locally sulphide-bearing and the Idefix sill (< 100 m) would not possess the heat required to effectively assimilate the country rocks. Although sulphidic country rock is not necessarily required for this deposit type, it remains a prospective characteristic if the intrusive sill is able to effectively extract crustal sulphide. The presence of stratiform pegmatitic gabbro with disseminated sulphide can be considered a good indicator of proximity to PGE-rich sulphide mineralisation in aphyric gabbro sills of the Montagnais Sill Complex. It is favourable that sills with stratiform gabbroic pegmatite are thick (> 100 m) and bound by other mafic-ultramafic sills, so that slower cooling rates may allow for pronged interaction between silicate melt and sulphide liquid.

# 7. Conclusions

The Idefix PGE-Cu prospect in the Labrador Trough, northern Québec represents a stack of differentiated gabbroic sills (~ 8 to 14 wt.% MgO, ~ 1 to 8 Cr/V), which are host to stratiform patchy net-textured and globular sulphide horizons, associated with stratiform gabbroic pegmatites. Primary minerals of the gabbroic hosts have undergone moderate to extensive alteration, whereby pyroxene is partially to completely replaced by amphibole, and plagioclase is almost completely saussurised. The patchy net-textured ores are more enriched

in chalcophile metals relative to the globular ores. Chalcophile metals in each horizon generally show good positive inter-element correlations (Ni/Cu = 0. 85; Cu/Pd = 0.68, Pd/Pt = 0.92, and IPGE/Ru = 0.96). Low Ni and IPGE tenors are ascribed to pre-emplacement fractionation of olivine and spinel. There is no evidence for the assimilation of crustal rock (La/Sm<sub>N</sub>  $\leq$  1 and Th/Yb<sub>N</sub>  $\leq$  1) or addition of exogenous sulphur (S/Se  $\leq$  4,00). Furthermore, *in situ* segregation of sulphide melt is precluded by insufficient volumes of PGE-undepleted magma to generate the observed PGE-Cu mineralisation. Henceforth, immiscible sulphide melt segregated before its final emplacement and was entrained upward. The patchy nettextured ores formed at *R* factors of up to 10,000, which was assisted by prolonged cooling rates due to thermal priming by antecedent pulses of magma. However, sills must be sufficiently thick ( $\geq$  100 m) and bound by hot mafic-ultramafic rocks to slow the cooling rate of the host sill. This allows for sulphide to interact with a greater volume of silicate magma and gives time to settle into narrow, PGE-rich reef-style horizons.

## **Funding**

W.D. Smith acknowledges the NERC GW4<sup>+</sup> doctoral training partnership (NE/L002434/1) for funding analytical procedures. Northern Shield Resources are thanked as acting as CASE partner for the project.

# Acknowledgements

The authors thank colleagues at Northern Shield Resources for fieldwork assistance, sampling, and data processing, as well as allowing publication of the data. Nick Giles is acknowledged for undertaking optical and electron microscopy of Idefix samples. Antony Oldroyd and Duncan Muir are all thanked for assistance and guidance during data

acquisition. This paper was greatly improved by thorough reviews from E. Mansur and C.M. 478 Lesher and additional comments from T.G. Blenkinsop. We thank Ali Polat for the editorial 479 handling of this manuscript. 480 481 References 482 Arndt, N. T., Czamanske, G. K., Walker, R. J., Chauvel, C., and Fedorenko, V. A. (2003). 483 Geochemistry and origin of the intrusive hosts of the Noril'sk-Talnakh Cu-Ni-PGE sulfide 484 485 deposits. Economic Geology, 98: 495-515. Avison, A. T. (1986). Report on 1986 Exploration Program in the Labrador Trough. 486 Geological Association of Canada. 487 Barnes, S-J., and Lightfoot, P.C. (2005). Formation of magmatic nickel-sulfide ore deposits 488 and processes affecting their copper and platinum-group element contents. In Hedenquist, 489 J.W., Thompson, J.F.H., Goldfarb, R.J. and Richards, J.P. Economic Geology 100: 179-213. 490 Barnes, S.-J., and Maier, W. D. (1999). The fractionation of Ni, Cu and the noble metals in 491 492 silicate and sulfide liquids. Geological Association of Canada, 13: 69–106. Barnes, S.-J., Picard, C., Giovenazzo, D., and Tremblay, C. (1992). The composition of 493 nickel-copper sulphide deposits and their host rocks from the Cape Smith Fold Belt, Northern 494 Quebec. Australian Journal of Earth Sciences, 39(3), 335-347. 495 Barnes, S. J., and Maier, W. D. (2002). Platinum-group elements and microstructures of 496 normal Merensky reef from Impala Platinum Mines, Bushveld Complex. Journal of 497 Petrology, 43(1), 103-128. 498 Barnes, S. J., & Liu, W. (2012). Pt and Pd mobility in hydrothermal fluids: evidence from 499

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## **Figure Captions**

Figure 1. a. Location of the study area in the lithotectonic divisions of Clark and Wares (2005). b. Geological map of the Idefix PGE-Cu prospect, showing the outline of the property and location of the boreholes addressed in this study. c. Cross-section across the centre of the property, showing the intersections of the labelled boreholes.

Figure 2. a. Schematic stratigraphy of the Idefix PGE-Cu prospect, showing the divisions of the Idefix sill. b. Textures and characteristics of the Baby Formation metasediments. c. Nature of the gabbroic pegmatite at the I1-I2 contact. d. Nature of the gabbroic pegmatite at the I3-PU contact. rill core is  $\sim$  4 cm in diameter. BF = Baby Formation, py = pyrite, qtz = quartz.

Figure 3. Texture and petrography of (a) the Primitive Unit. (b) Idefix unit I3 and I2, (c) Idefix unit I1. d. CIPW normative mineralogy of gabbroic and gabbroic pegmatitic rocks at Idefix. cpx = clinopyroxene, plg = plagioclase, amp = amphibole, qtz = quartz.

Figure 4. a. Texture and mineralogy of patchy net-textured sulphide. b. Texture and mineralogy of globular sulphide. Note the elliptical shape of the sulphide globules. po = pyrrhotite, pn = pentlandite, ccp = chalcopyrite, ars = sulpharsenides.

Figure 5. a-f. MgO against CaO, FeOt, TiO<sub>2</sub>, Cr/V, Sr, and Ni.

Figure 6. a-b. Primitive mantle normalised (Sun and McDonough 1989) lithophile multielement plots.

Figure 7. a-c. Sulphur against Ni, Cu, and 2PGE + Au. d. Pd against Pt, whereby net-textured and globular ores correlate at Pt/Pd values of ~ 2.8. e. Cu against 2PGE + Au. f. Ni against 2PGE + Au. Note the different trends in net-textured and globular ores in plots c, e, and f.

Figure 8. a-f. Ir against Ni, Ru, Rh, Pd, Pt, and Au. Note that IPGE and Pt show good positive correlations ( $R^2 > 0.8$ ) and that all samples plot with Pd/Ir values below 500. RG = regional gabbro.

Figure 9. Primitive mantle normalised (Barnes and Maier 1999) chalcophile multi-element plots for (a) patchy net-textured ores and (b) globular ores. For comparison, profiles form the J-M Reef (Godel *et al.* 2002), Roby Zone of Lac des Iles (LDI; Hinchey *et al.* 2005), Merensky Reef (Barnes and Maier 2002), the Platreef at Rooipoort (Maier *et al.* 2008a), and the Paladin deposit in the Labrador Trough (Clark and Wares 2005). All samples with > 0.5 wt.% S have been normalised to 100% sulphide using the method of Barnes and Lightfoot (2005).

Figure 10. Pd against Cu/Pd with marginal histograms. The underlain grey field represents the expected Cu/Pd range of mantle rock (Barnes and Maier 1999). Grey boxes represent the composition of sulphide at different whole-rock volumes at different *R* factors (see text for discussion).

Figure 11. Downhole geochemistry of borehole 13ID-13, showing the downward trend of lithophile and chalcophile elements.

Figure 12. a-b. La/Sm<sub>N</sub> against La/Yb<sub>N</sub> and La/Nb<sub>N</sub> against Th/Yb<sub>N</sub>, overlain with binary mixing models between Hellancourt basalt (Ciborowski *et al.* 2017) and Baby Formation sediments. c. S against Se underlain with that of mantle range (2850 to 4300; Eckstrand and Hulbert 1987). Note that all samples plot at or just below that expected for mantle rock. d. S/Se against La/Sm<sub>CN</sub> showing no changes in La/Sm<sub>N</sub> values relative to S/Se. Normalised ratios were normalised using values of Sun and McDonough (1989).

Figure 13. Emplacement scenarios for the gabbroic rocks at Idefix if (1) sequentially emplaced and overturned, (2) non-sequentially emplaced and overturned, (3) sequentially emplaced, and (4) non-sequentially emplaced.

 Table 1. Representative whole-rock compositions of the rocks at Idefix. Full data reported in Supplementary data 2.

Rock Type	PU	PU	Transitional	I3	I3	I2	I2	I1	I1	Sediments
Hole ID	13ID-13	13ID-13	13ID-04	13ID-13	13ID-02	13ID-02	13ID-04	13ID-13	13ID-02	13ID-07
Depth (m)	15.2	62	18.6	116.6	41	88.8	69.7	191.7	118.6	81
Major Elements (wt.%)										
$SiO_2$	49.5	49.3	49.6	48.5	48.5	48.4	48.2	48.6	48.5	59.8
$TiO_2$	0.3	0.3	0.4	0.4	0.4	0.5	0.5	0.5	0.6	0.4
$Al_2O_3$	11.5	13.1	13.7	13.1	13.6	14.6	14.6	13.6	14.0	14.2
$Fe_2O_3$	7.2	7.7	8.0	8.5	8.9	9.5	9.6	9.5	10.7	11.6
MnO	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2
MgO	11.6	11.2	10.7	10.9	11.1	10.0	10.2	9.6	9.7	2.9
CaO	16.1	14.5	15.0	13.8	13.9	12.9	13.3	13.5	12.3	2.7
$Na_2O$	0.6	0.9	1.1	1.1	1.1	1.2	1.3	1.1	1.1	2.6
$K_2O$	0.0	0.0	0.1	0.1	0.1	0.1	0.1	0.1	0.4	3.8
$P_2O_5$	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.1
LOI	2.1	2.2	2.2	2.3	2.3	2.6	2.7	2.1	0.5	1.2
Total	99.3	99.6	100.9	98.9	100.0	100.0	100.6	98.9	98.0	99.6
Trace Elemen	ts (ppm)									
S (wt.%)	0.0	0.0	0.0	0.1	0.1	0.1	0.1	0.1	0.4	0.1
Sc	52	44	45	42	42	36	36	38	32	9
V	194	195	204	181	199	214	210	229	226	53
Cr	1890	1400	1390	1120	1250	800	840	880	710	50
Co	38	42	37	46	48	47	42	47	54	11
Ni	171	159	137	261	256	179	175	171	331	22
Cu	61	58	55	484	381	195	128	207	710	5
Zn	50	43	43	56	52	58	59	62	67	111
As	1.8	0.6	0.7	0.6	1.1	8.5	1.6	19.0	41.1	0.5
Rb	0.2	1.0	0.8	1.2	1.7	3.7	1.4	1.4	11.6	162.0
Sr	54	65	74	62	66	74	73	84	83	219
Y	7.6	7.9	9.6	8.5	9.1	10.4	10.3	11.7	11.5	14.8
Zr	7	10	17	13	17	21	22	28	29	174
Nb	0.6	0.7	0.7	0.8	0.8	1.0	1.1	1.4	1.3	7.6
Sb	0.1	0.1	0.1	0.2	0.1	0.2	0.3	0.2	0.2	0.1
Cs	< 0.01	0.0	0.0	0.0	0.1	0.1	0.0	0.0	0.1	7.6
Ba	5	10	10	12	15	22	10	10	48	426
Th	0.1	0.1	0.2	0.1	0.1	0.2	0.2	0.2	0.2	13.5
Precious Meta										
Pd	bdl	0.02	0.01	0.25	0.21	0.06	0.05	0.06	0.17	bdl
Pt	bdl	0.02	0.01	0.09	0.07	0.03	0.02	0.02	0.05	bdl
Au	bdl	bdl	bdl	0.02	0.02	bdl	bdl	bdl	0.01	bdl
REEs (ppm)										
La	1.3	0.9	2.4	1.0	1.2	1.5	1.5	1.7	1.7	43.9
Ce	2.0	2.3	4.4	2.7	3.2	3.9	3.8	4.0	4.6	79.1
Pr	0.3	0.4	0.6	0.5	0.5	0.7	0.6	0.7	0.7	8.6
Nd	1.8	2.0	3.0	2.4	2.7	3.0	3.0	3.9	3.6	30.5
Sm	0.6	0.8	0.9	0.9	0.8	1.1	1.1	1.2	1.3	4.8

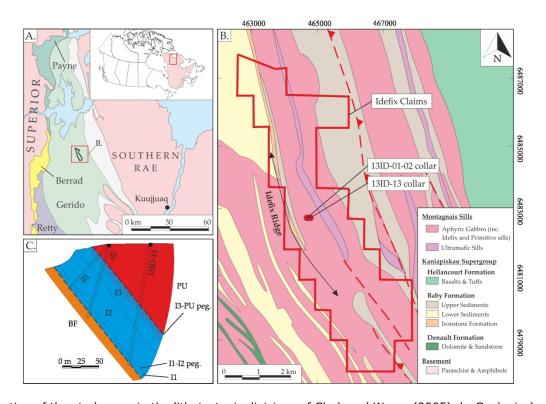
Eu         0.3         0.3         0.4         0.4         0.4         0.4         0.5         0.5         0.5         1.0           Gd         1.0         1.1         1.5         1.4         1.4         1.6         1.7         1.8         1.6         3.9           Tb         0.2         0.2         0.2         0.2         0.2         0.2         0.2         0.3         0.3         0.3         0.5           Dy         1.5         1.4         1.6         1.5         1.6         1.7         1.8         1.9         2.0         3.0           Ho         0.3         0.3         0.3         0.4         0.3         0.4         0.5         0.4         0.6           Er         0.8         0.8         1.1         1.0         1.1         1.1         1.1         1.4         1.2         1.7           Tm         0.1         0.1         0.1         0.2         0.1         0.1         0.2         0.2         0.2         0.2         0.2           Yb         0.9         0.9         1.0         1.0         1.0         1.2         1.1         1.3         1.2         1.2           Lu											
Tb         0.2         0.2         0.2         0.2         0.2         0.2         0.2         0.3         0.3         0.3         0.5           Dy         1.5         1.4         1.6         1.5         1.6         1.7         1.8         1.9         2.0         3.0           Ho         0.3         0.3         0.4         0.3         0.4         0.4         0.5         0.4         0.6           Er         0.8         0.8         1.1         1.0         1.1         1.1         1.1         1.4         1.2         1.7           Tm         0.1         0.1         0.1         0.2         0.1         0.1         0.2         0.2         0.2         0.2         0.2           Yb         0.9         0.9         1.0         1.0         1.0         1.2         1.1         1.3         1.2         1.2           Lu         0.1         0.1         0.1         0.1         0.2         0.2         0.2         0.2         0.2           Element Ratios           Mg#         58.4         54.8         54.1         52.8         56.1         51.4         48.3         46.9         49.8 <t< td=""><td>Eu</td><td>0.3</td><td>0.3</td><td>0.4</td><td>0.4</td><td>0.4</td><td>0.4</td><td>0.5</td><td>0.5</td><td>0.5</td><td>1.0</td></t<>	Eu	0.3	0.3	0.4	0.4	0.4	0.4	0.5	0.5	0.5	1.0
Dy         1.5         1.4         1.6         1.5         1.6         1.7         1.8         1.9         2.0         3.0           Ho         0.3         0.3         0.3         0.4         0.3         0.4         0.5         0.4         0.6           Er         0.8         0.8         1.1         1.0         1.1         1.1         1.1         1.4         1.2         1.7           Tm         0.1         0.1         0.1         0.2         0.1         0.1         0.2         0.2         0.2         0.2         0.2           Yb         0.9         0.9         1.0         1.0         1.0         1.2         1.1         1.3         1.2         1.2           Lu         0.1         0.1         0.1         0.1         0.1         0.2         0.2         0.2         0.2         0.2         0.2           Element Ratios           Mg#         58.4         54.8         54.1         52.8         56.1         51.4         48.3         46.9         49.8         20.0           Cr/V         9.7         7.2         6.8         6.2         6.3         3.7         4.0         3.8	Gd	1.0	1.1	1.5	1.4	1.4	1.6	1.7	1.8	1.6	3.9
Ho 0.3 0.3 0.3 0.4 0.3 0.4 0.4 0.5 0.4 0.6  Er 0.8 0.8 1.1 1.0 1.1 1.1 1.1 1.1 1.4 1.2 1.7  Tm 0.1 0.1 0.1 0.1 0.2 0.1 0.1 0.2 0.2 0.2 0.2 0.2  Yb 0.9 0.9 1.0 1.0 1.0 1.0 1.2 1.1 1.3 1.2 1.2  Lu 0.1 0.1 0.1 0.1 0.1 0.1 0.2 0.2 0.2 0.2 0.2  Element Ratios  Mg# 58.4 54.8 54.1 52.8 56.1 51.4 48.3 46.9 49.8 20.0  Cr/V 9.7 7.2 6.8 6.2 6.3 3.7 4.0 3.8 3.1 0.9  Eu/Eu* 1.2 0.9 0.9 1.0 1.0 0.9 1.0 0.9 1.0 0.9 1.0 0.7  Ni/Cu 2.8 2.7 2.5 0.5 0.7 0.9 1.4 0.8 0.5 4.4  Cu/Pd 15250 2900 4583 1928 1841 3095 2667 3632 4251 5000	Tb	0.2	0.2	0.2	0.2	0.2	0.2	0.3	0.3	0.3	0.5
Er         0.8         0.8         1.1         1.0         1.1         1.1         1.1         1.4         1.2         1.7           Tm         0.1         0.1         0.1         0.2         0.1         0.1         0.2         0.2         0.2         0.2         0.2           Yb         0.9         0.9         1.0         1.0         1.0         1.2         1.1         1.3         1.2         1.2           Lu         0.1         0.1         0.1         0.1         0.1         0.2         0.2         0.2         0.2         0.2         0.2           Element Ratios           Mg#         58.4         54.8         54.1         52.8         56.1         51.4         48.3         46.9         49.8         20.0           Cr/V         9.7         7.2         6.8         6.2         6.3         3.7         4.0         3.8         3.1         0.9           Eu/Eu*         1.2         0.9         0.9         1.0         1.0         0.9         1.0         0.9         1.0         0.7           Ni/Cu         2.8         2.7         2.5         0.5         0.7         0.9         1.4	Dy	1.5	1.4	1.6	1.5	1.6	1.7	1.8	1.9	2.0	3.0
Tm         0.1         0.1         0.1         0.2         0.1         0.1         0.2         0.2         0.2         0.2         0.2         0.2           Yb         0.9         0.9         1.0         1.0         1.0         1.2         1.1         1.3         1.2         1.2           Lu         0.1         0.1         0.1         0.1         0.1         0.2         0.2         0.2         0.2         0.2           Element Ratios           Mg#         58.4         54.8         54.1         52.8         56.1         51.4         48.3         46.9         49.8         20.0           Cr/V         9.7         7.2         6.8         6.2         6.3         3.7         4.0         3.8         3.1         0.9           Eu/Eu*         1.2         0.9         0.9         1.0         1.0         0.9         1.0         0.9         1.0         0.7           Ni/Cu         2.8         2.7         2.5         0.5         0.7         0.9         1.4         0.8         0.5         4.4           Cu/Pd         15250         2900         4583         1928         1841         3095 <td< td=""><td>Но</td><td>0.3</td><td>0.3</td><td>0.3</td><td>0.4</td><td>0.3</td><td>0.4</td><td>0.4</td><td>0.5</td><td>0.4</td><td>0.6</td></td<>	Но	0.3	0.3	0.3	0.4	0.3	0.4	0.4	0.5	0.4	0.6
Yb         0.9         0.9         1.0         1.0         1.0         1.2         1.1         1.3         1.2         1.2           Lu         0.1         0.1         0.1         0.1         0.2         0.2         0.2         0.2         0.2         0.2           Element Ratios           Mg#         58.4         54.8         54.1         52.8         56.1         51.4         48.3         46.9         49.8         20.0           Cr/V         9.7         7.2         6.8         6.2         6.3         3.7         4.0         3.8         3.1         0.9           Eu/Eu*         1.2         0.9         0.9         1.0         1.0         0.9         1.0         0.9         1.0         0.7           Ni/Cu         2.8         2.7         2.5         0.5         0.7         0.9         1.4         0.8         0.5         4.4           Cu/Pd         15250         2900         4583         1928         1841         3095         2667         3632         4251         5000	Er	0.8	0.8	1.1	1.0	1.1	1.1	1.1	1.4	1.2	1.7
Lu         0.1         0.1         0.1         0.1         0.1         0.2         0.2         0.2         0.2         0.2         0.2           Element Ratios           Mg#         58.4         54.8         54.1         52.8         56.1         51.4         48.3         46.9         49.8         20.0           Cr/V         9.7         7.2         6.8         6.2         6.3         3.7         4.0         3.8         3.1         0.9           Eu/Eu*         1.2         0.9         0.9         1.0         1.0         0.9         1.0         0.9         1.0         0.7           Ni/Cu         2.8         2.7         2.5         0.5         0.7         0.9         1.4         0.8         0.5         4.4           Cu/Pd         15250         2900         4583         1928         1841         3095         2667         3632         4251         5000	Tm	0.1	0.1	0.1	0.2	0.1	0.1	0.2	0.2	0.2	0.2
Element Ratios         Mg#       58.4       54.8       54.1       52.8       56.1       51.4       48.3       46.9       49.8       20.0         Cr/V       9.7       7.2       6.8       6.2       6.3       3.7       4.0       3.8       3.1       0.9         Eu/Eu*       1.2       0.9       0.9       1.0       1.0       0.9       1.0       0.9       1.0       0.7         Ni/Cu       2.8       2.7       2.5       0.5       0.7       0.9       1.4       0.8       0.5       4.4         Cu/Pd       15250       2900       4583       1928       1841       3095       2667       3632       4251       5000	Yb	0.9	0.9	1.0	1.0	1.0	1.2	1.1	1.3	1.2	1.2
Mg#         58.4         54.8         54.1         52.8         56.1         51.4         48.3         46.9         49.8         20.0           Cr/V         9.7         7.2         6.8         6.2         6.3         3.7         4.0         3.8         3.1         0.9           Eu/Eu*         1.2         0.9         0.9         1.0         0.9         1.0         0.9         1.0         0.7           Ni/Cu         2.8         2.7         2.5         0.5         0.7         0.9         1.4         0.8         0.5         4.4           Cu/Pd         15250         2900         4583         1928         1841         3095         2667         3632         4251         5000	Lu	0.1	0.1	0.1	0.1	0.1	0.2	0.2	0.2	0.2	0.2
Ct/V         9.7         7.2         6.8         6.2         6.3         3.7         4.0         3.8         3.1         0.9           Eu/Eu*         1.2         0.9         0.9         1.0         1.0         0.9         1.0         0.9         1.0         0.7           Ni/Cu         2.8         2.7         2.5         0.5         0.7         0.9         1.4         0.8         0.5         4.4           Cu/Pd         15250         2900         4583         1928         1841         3095         2667         3632         4251         5000	Element Rati	os									
Eu/Eu*       1.2       0.9       0.9       1.0       1.0       0.9       1.0       0.9       1.0       0.7         Ni/Cu       2.8       2.7       2.5       0.5       0.7       0.9       1.4       0.8       0.5       4.4         Cu/Pd       15250       2900       4583       1928       1841       3095       2667       3632       4251       5000	Mg#	58.4	54.8	54.1	52.8	56.1	51.4	48.3	46.9	49.8	20.0
Ni/Cu 2.8 2.7 2.5 0.5 0.7 0.9 1.4 0.8 0.5 4.4 Cu/Pd 15250 2900 4583 1928 1841 3095 2667 3632 4251 5000	Cr/V	9.7	7.2	6.8	6.2	6.3	3.7	4.0	3.8	3.1	0.9
Cu/Pd 15250 2900 4583 1928 1841 3095 2667 3632 4251 5000	Eu/Eu*	1.2	0.9	0.9	1.0	1.0	0.9	1.0	0.9	1.0	0.7
	Ni/Cu	2.8	2.7	2.5	0.5	0.7	0.9	1.4	0.8	0.5	4.4
Pd/Pt 1.0 1.0 1.0 2.7 2.9 2.2 2.3 2.9 3.1 1.3	Cu/Pd	15250	2900	4583	1928	1841	3095	2667	3632	4251	5000
	Pd/Pt	1.0	1.0	1.0	2.7	2.9	2.2	2.3	2.9	3.1	1.3

Eu/Eu\* calculated by Eu $_{\! N}\,/\,(Sm_{\! N}$  \*  $Gd_{\! N})^{0.5}$ 

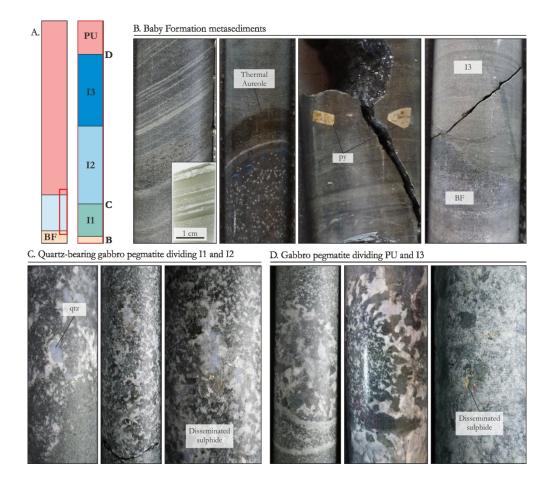
 Table 2. Chalcophile element concentrations of gabbroic rocks at Idefix.

Sample	Unit		Base Metal (wt.%)		Platinum Group Elements & Gold (ppb)*						
		Ni	Cu	Ir	Ru	Rh	Pt	Pd	Au	(wt.%)	
1D11-02	I1	2.1	5.2	55	69	642	2166	17228	173	2.0	
1D11-07D	I1	2.9	8.7	118	88	1086	14650	64878	1036	0.8	
1D11-14	I1	2.4	2.3	117	95	859	6361	18375	348	2.3	
1D11-04	I3	3.2	14.0	372	211	3119	20354	68939	3578	2.8	
1D11-06	I3	3.2	11.3	328	221	2968	22333	74725	2486	0.8	
1D11-12	I3	0.4	26.7	331	178	2887	20619	67258	12670	3.2	
1D11-13	I3	3.1	19.0	827	477	6794	43641	156154	3205	1.3	
1D11-24	I3	0.7	14.1	725	415	6080	2372	76511	1732	1.6	
1D11-01	Barren	2.1	8.2	330	194	2969	30544	100081	2008	0.5	

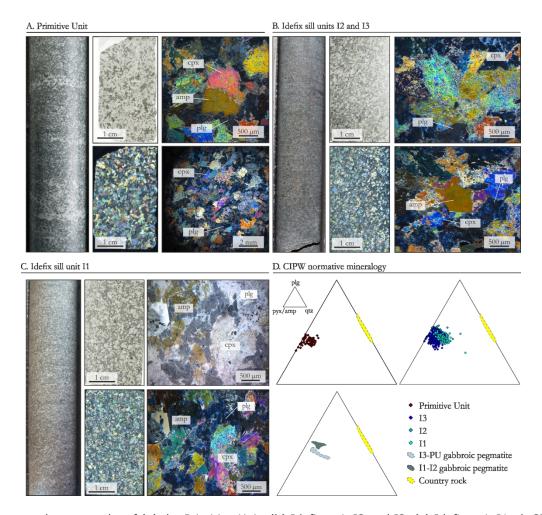
<sup>\*</sup>Normalised to 100% sulphide using the method of Barnes & Lightfoot (2005).



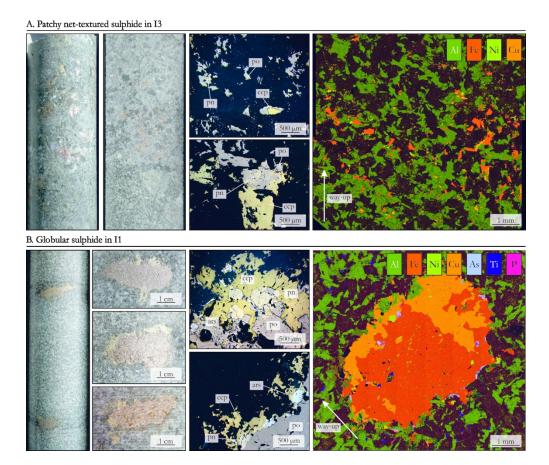
a. Location of the study area in the lithotectonic divisions of Clark and Wares (2005). b. Geological map of the Idefix PGE-Cu prospect, showing the outline of the property and location of the boreholes addressed in this study. c. Cross-section across the centre of the property, showing the intersections of the labelled boreholes.



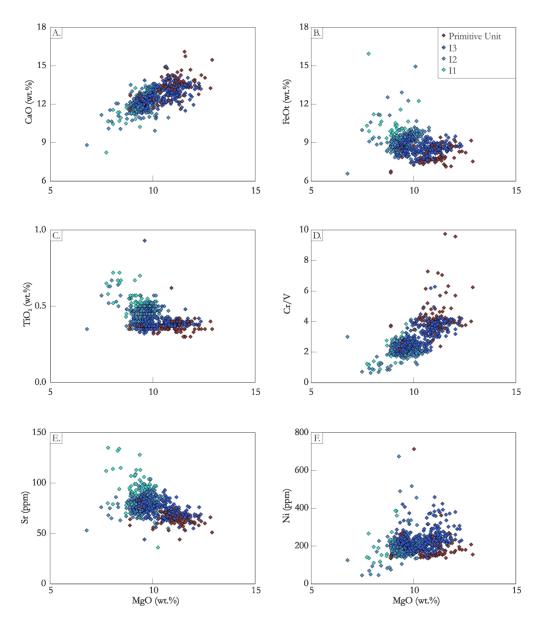
a. Schematic stratigraphy of the Idefix PGE-Cu prospect, showing the divisions of the Idefix sill. b. Textures and characteristics of the Baby Formation metasediments. c. Nature of the gabbroic pegmatite at the I1-I2 contact. d. Nature of the gabbroic pegmatite at the I3-PU contact. rill core is  $\sim 4$  cm in diameter. BF = Baby Formation, py = pyrite, qtz = quartz.



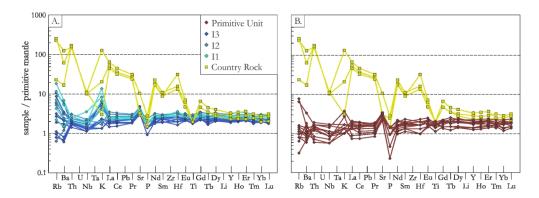
Texture and petrography of (a) the Primitive Unit. (b) Idefix unit I3 and I2, (c) Idefix unit I1. d. CIPW normative mineralogy of gabbroic and gabbroic pegmatitic rocks at Idefix. cpx = clinopyroxene, plg = plagioclase, amp = amphibole, qtz = quartz.



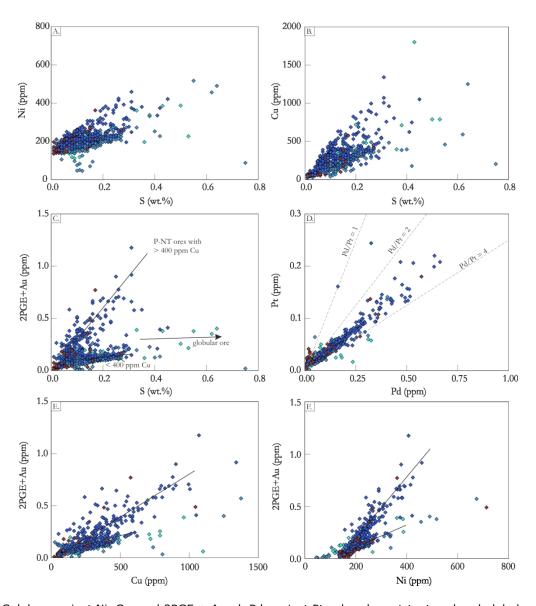
a. Texture and mineralogy of patchy net-textured sulphide. b. Texture and mineralogy of globular sulphide. Note the elliptical shape of the sulphide globules. po = pyrrhotite, po = pyrrhotite, po = pentlandite, po = pyrrhotite, po



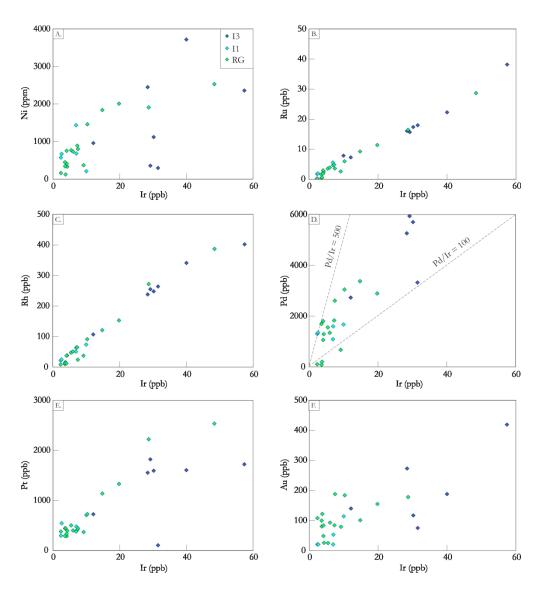
a-f. MgO against CaO, FeOt, TiO2, Cr/V, Sr, and Ni.



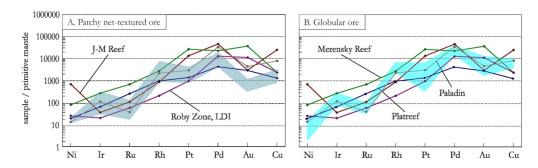
a-b. Primitive mantle normalised (Sun and McDonough 1989) lithophile multi-element plots.



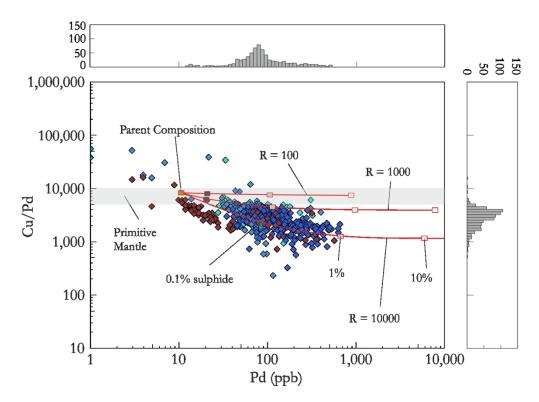
a-c. Sulphur against Ni, Cu, and 2PGE + Au. d. Pd against Pt, whereby net-textured and globular ores correlate at Pt/Pd values of  $\sim$  2.8. e. Cu against 2PGE + Au. f. Ni against 2PGE + Au. Note the different trends in net-textured and globular ores in plots c, e, and f.



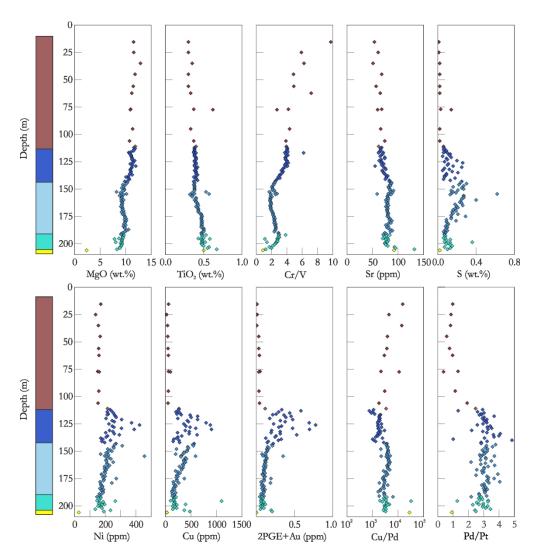
a-f. Ir against Ni, Ru, Rh, Pd, Pt, and Au. Note that IPGE and Pt show good positive correlations (R2 > 0.8) and that all samples plot with Pd/Ir values below 500. RG = regional gabbro.



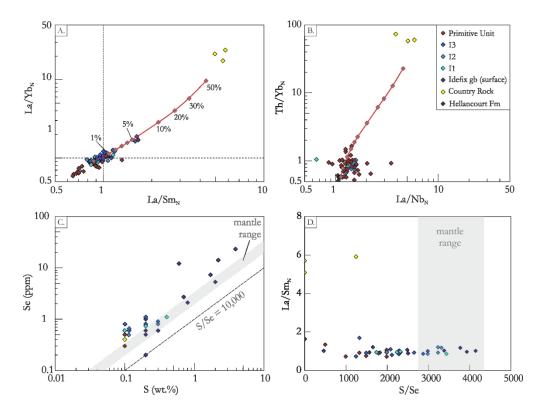
Primitive mantle normalised (Barnes and Maier 1999) chalcophile multi-element plots for (a) patchy net-textured ores and (b) globular ores. For comparison, profiles form the J-M Reef (Godel et al. 2002), Roby Zone of Lac des Iles (LDI; Hinchey et al. 2005), Merensky Reef (Barnes and Maier 2002), the Platreef at Rooipoort (Maier et al. 2008a), and the Paladin deposit in the Labrador Trough (Clark and Wares 2005). All samples with > 0.5 wt.% S have been normalised to 100% sulphide using the method of Barnes and Lightfoot (2005).



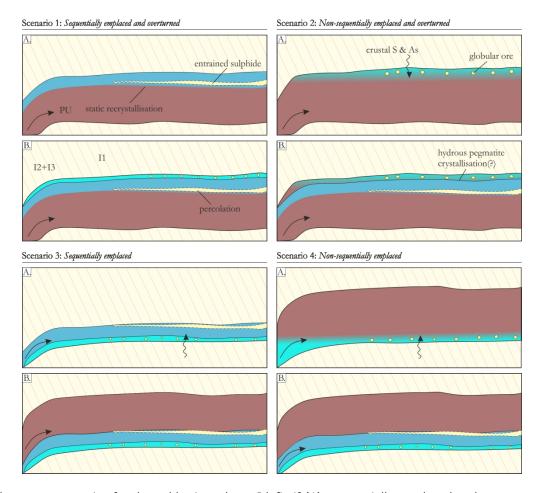
Pd against Cu/Pd with marginal histograms. The underlain grey field represents the expected Cu/Pd range of mantle rock (Barnes and Maier 1999). Grey boxes represent the composition of sulphide at different whole-rock volumes at different R factors (see text for discussion).



Downhole geochemistry of borehole 13ID-13, showing the downward trend of lithophile and chalcophile elements.



a-b. La/SmN against La/YbN and La/NbN against Th/YbN, overlain with binary mixing models between Hellancourt basalt (Ciborowski et al. 2017) and Baby Formation sediments. c. S against Se underlain with that of mantle range (2850 to 4300; Eckstrand and Hulbert 1987). Note that all samples plot at or just below that expected for mantle rock. d. S/Se against La/SmCN showing no changes in La/SmN values relative to S/Se. Normalised ratios were normalised using values of Sun and McDonough (1989).



Emplacement scenarios for the gabbroic rocks at Idefix if (1) sequentially emplaced and overturned, (2) non-sequentially emplaced and overturned, (3) sequentially emplaced, and (4) non-sequentially emplaced.