



## Article

# Ferrodymolybdenite, $\text{FeMo}^{3+}_2\text{S}_4$ from Daba-Siwaqa, Jordan – the first natural compound of trivalent molybdenum

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## Abstract

Ferrodymolybdenite with ideal formula  $\text{FeMo}^{3+}_2\text{S}_4$  ( $C2/c$ ,  $a = 11.8249(8) \text{ \AA}$ ,  $b = 6.5534(3) \text{ \AA}$ ,  $c = 13.0052(10) \text{ \AA}$ ,  $\beta = 114.474(9)^\circ$ ,  $V = 917.27(12) \text{ \AA}^3$  and  $Z = 8$ ) was discovered in a differentiated sulfide nodule composed of troilite and pentlandite parts. The nodule was detected in the central zone of a diopside–anorthite–tridymite oval paralava body, ~30 metres in diameter, within the pyrometamorphic Hatrurim Complex in Daba-Siwaqa, Jordan. Ferrodymolybdenite is the first trivalent molybdenum compound discovered in Nature. Its synthetic analogue crystallises in the  $C1c1$  space group. Ferrodymolybdenite with the empirical formula  $(\text{Fe}^{2+}_{0.99}\text{Cu}^{2+}_{0.07}\text{Ni}^{2+}_{0.04})_{\Sigma 1.10}\text{Mo}^{3+}_{1.94}(\text{S}^{2-}_{3.98}\text{P}^{3-}_{0.02})_{\Sigma 4.00}$  was identified in the troilite part of the differentiated sulfide nodule. The nodule contains inclusions of tetrataenite, nickelporphide, molybdenite, galena and rudashevskyite. Ferrodymolybdenite forms platy crystals with dimensions ranging from  $3 \times 100 \mu\text{m}$  to  $20 \times 40 \mu\text{m}$ . The mineral exhibits a grey colour and a dark grey streak. It is opaque with a metallic lustre, and its Mohs hardness is ~3. The cleavage observed in the mineral is perfect on  $\{001\}$ , good on  $\{100\}$  and poor on  $\{010\}$ . Its tenacity is sectile, and its fracture is smooth. The calculated density of  $5.445 \text{ g}\cdot\text{cm}^{-3}$  was derived from the empirical formula and unit cell volume refined from single-crystal X-ray diffraction data. In reflected light, ferrodymolybdenite appears grey to light grey with a blueish tinge. It is anisotropic, with a reflectance in the range of 34–40%. The crystallisation of ferrodymolybdenite occurred in reduced conditions in monosulfide  $\text{Fe}(\text{+Ni})$  melt at a temperature of 1000–1100°C and at low pressure.

**Keywords:** new mineral; ferrodymolybdenite; crystal structure;  $\text{Mo}^{3+}$ ; Daba-Siwaqa; Jordan; Hatrurim Complex

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## Introduction

Ferrodymolybdenite,  $\text{FeMo}^{3+}_2\text{S}_4$  ( $C2/c$ ,  $a = 11.8249(8) \text{ \AA}$ ,  $b = 6.5534(3) \text{ \AA}$ ,  $c = 13.0052(10) \text{ \AA}$ ,  $\beta = 114.474(9)^\circ$ ,  $V = 917.27(12) \text{ \AA}^3$  and  $Z = 8$ ) was identified in a sulfide nodule (~7 mm) in the central part of an oval diopside–anorthite–tridymite paralava body with a diameter of ~30 m within a field of spurrite-bearing rocks of the Hatrurim Complex, Daba-Siwaqa, Jordan. This paralava is the site of the first description of 11 new phosphides and phosphates, which were found to be concentrated along the contact between the paralava and host altered sedimentary rocks (Britvin *et al.*, 2015; Galuskin *et al.*, 2025). The mineral composition and conditions of paralava formation, as well as the genetic

aspects of phosphide and phosphate crystallisation in this paralava, are considered elsewhere (Galuskin *et al.*, 2023a, 2025).

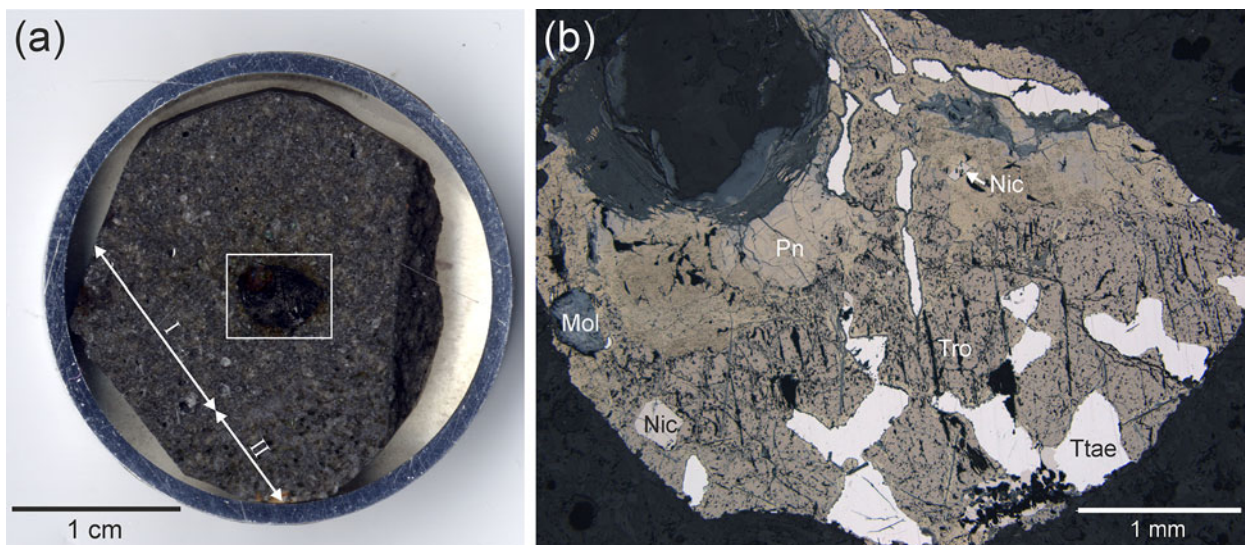
Ferrodymolybdenite is the first natural compound of trivalent molybdenum. All known minerals contain molybdenum in an oxidation state of +4 or +6; however, molybdenum can be found in a reduced form in phosphides (Murashko *et al.*, 2022). A metallic Mo, hexamolybdenum, has been identified in the Allende meteorite (Ma *et al.*, 2014). It has previously been determined that the synthetic compound  $\text{FeMo}^{3+}_2\text{S}_4$  belongs to the  $C1c1$  space group ( $a = 11.8148(2) \text{ \AA}$ ,  $b = 6.5499(1) \text{ \AA}$ ,  $c = 13.014(2) \text{ \AA}$ ,  $\beta = 114.455(1)^\circ$  and  $V = 916.75 \text{ \AA}^3$ ) (Guillevis *et al.*, 1974; Vaqueiro *et al.*, 2002), whereas for ferrodymolybdenite we have accepted a model with the  $C2/c$  space group.

The present article provides a description of a new mineral, ferrodymolybdenite,  $\text{FeMo}^{3+}_2\text{S}_4$ . The mineral and name have been approved by the Commission on New Minerals, Nomenclature and Classification (CNMNC) of the International Mineralogical Association (IMA), IMA2023–019 (mineral symbol – Fdmol). The mineral name ferrodymolybdenite is derived from its composition

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**Figure 1.** (a) Sulfide nodules in diopside-anorthite-tridymite paralava. I - porous rock and II - massive rock. The fragment outlined (holotype specimen 6005/1) is enlarged in Fig. 1b. (b) Reflected light image of the differentiated sulfide nodule with a gas bubble (upper part) and composed of troilite and pentlandite zones. Relatively large inclusions of tetrataenite and nickelporphide are present in the troilite zone. Tro = troilite; Ttae = tetrataenite; Mol = molybdenite; Pn = pentlandite and Nic = nickelporphide [Mineral symbols here and below according to Warr (2021)].

and structure: ‘ferro’ refers to  $\text{Fe}^{2+}$ , whereas ‘di+molybdenite’ indicates the presence of two  $\text{MoS}_2$  layers. Type material has been deposited in the mineralogical collection of the Fersman Mineralogical Museum, Leninsky pr., 18/k. 2, 115162 Moscow, Russia, registration number: 6005/1.

## Methods of investigation

The morphology and composition of ferrodymolybdenite and associated minerals were studied using optical microscopy, scanning electron microscopes (Phenom XL and Quanta 250, Institute of Earth Sciences, Faculty of Natural Sciences, University of Silesia, Sosnowiec, Poland) and an electron microprobe analyser (Cameca SX100, Micro-Area Analysis Laboratory, Polish Geological Institute – National Research Institute, Warsaw, Poland). Chemical analyses of sulfide nodule minerals were conducted in WDS mode (wavelength-dispersive spectroscopy, settings: 15 kV, 20 nA and  $\sim 1 \mu\text{m}$  beam diameter) using the following lines and standards:  $\text{MoL}\beta$  – Mo;  $\text{CuK}\alpha$  – Cu;  $\text{NiK}\alpha$  – pentlandite;  $\text{FeK}\alpha$  –  $\text{FeS}_2$ ;  $\text{SK}\alpha$  –  $\text{FeS}_2$ ;  $\text{ZnS}$ ;  $\text{PK}\alpha$  – InP;  $\text{SiK}\alpha$ ,  $\text{MgK}\alpha$  – diopside;  $\text{AlK}\alpha$  – orthoclase;  $\text{SeL}\beta$  –  $\text{In}_2\text{Se}_3$ ;  $\text{AsL}\beta$ ,  $\text{CoK}\alpha$  –  $\text{CoNiAs}_3$ ;  $\text{CaK}\alpha$  – apatite;  $\text{SbLa}$  –  $\text{Sb}_2\text{Te}_3$ ;  $\text{PbM}\beta$  – pyromorphite;  $\text{ZnK}\alpha$  –  $\text{ZnS}$ ;  $\text{MnK}\alpha$  – rhodonite;  $\text{CrK}\alpha$  –  $\text{Cr}_2\text{O}_3$ ;  $\text{VK}\alpha$  – V; and  $\text{TiK}\alpha$  – rutile. Other chemical elements were below the detection limit.

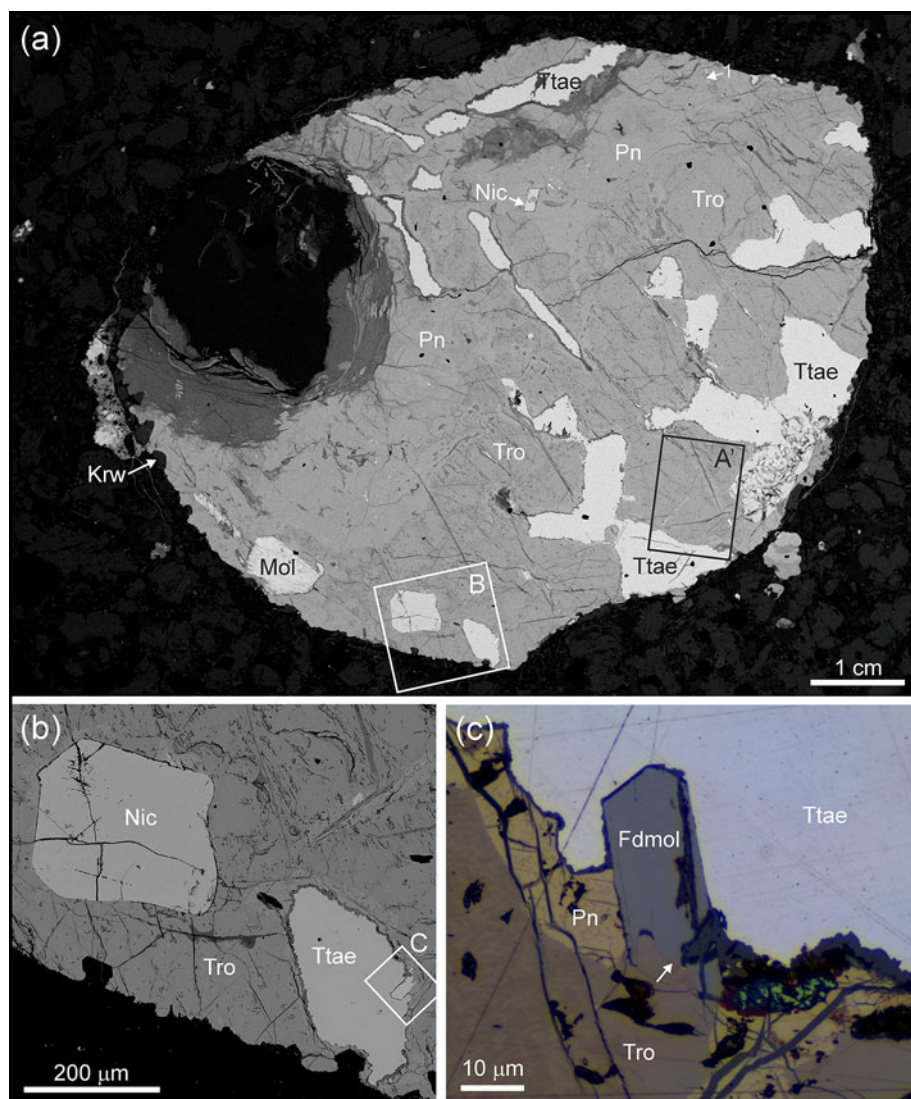
The Raman spectra of ferrodymolybdenite and molybdenite were recorded on a WITec alpha 300R Confocal Raman Microscope (Department of Earth Science, University of Silesia, Poland) equipped with an air-cooled solid laser (488 nm) and a CCD camera operating at  $-61^\circ\text{C}$ . The laser radiation was coupled to a microscope through a single-mode optical fibre with a diameter of  $3.5 \mu\text{m}$ . An air Zeiss LD EC Epiplan-Neofluar DIC-100/0.75NA objective was used. The Raman scattered light was focused onto a multi-mode fibre and monochromator with a  $1800 \text{ gr mm}^{-1}$  grating. The laser power at the sample position was 5–7 mW. Twenty scans with an integration time of 3 s and a resolution of  $2 \text{ cm}^{-1}$  were collected and averaged. The spectrometer

monochromator was calibrated using the Raman scattering line of a silicon plate ( $520.7 \text{ cm}^{-1}$ ).

Single-crystal X-ray diffraction (XRD) studies of ferrodymolybdenite were carried out using a SuperNova diffractometer with a mirror monochromator ( $\text{MoK}\alpha$  and  $\lambda = 0.71073 \text{ \AA}$ ) and an Atlas CCD detector (formerly Agilent Technologies, currently Rigaku Oxford Diffraction) at the Institute of Physics, University of Silesia, Poland. Single-crystal XRD data were collected using a ferrodymolybdenite crystal fragment  $40 \times 30 \times 10 \mu\text{m}$  in size. The ferrodymolybdenite structure was refined using the *SHELX-2019/2* program (Sheldrick, 2015). The crystal structure was refined starting from the atomic coordinates of synthetic  $\text{FeMo}_2\text{S}_4$  (Vaqueiro *et al.*, 2002).

## Background information

The pyrometamorphic rocks of the Hatrurim Complex, represented by spurrite, larnite and gehlenite rocks, form large fields along the Dead Sea Rift in Israel, Palestine and Jordan. Descriptions of the Hatrurim Complex can be found in numerous publications (Bentor, 1960; Gross, 1977; Vapnik *et al.*, 2007; Geller *et al.*, 2012; Novikov *et al.*, 2013; Khoury *et al.*, 2014; Galuskina *et al.*, 2014). The genesis of the Complex remains an unsolved problem, but all researchers agree that the rocks of the Hatrurim Complex are products of combustion metamorphism. Two genetic hypotheses have been proposed, the first of which relates pyrometamorphic transformations of the sedimentary protolith to the combustion of dispersed organic fuels (Kolodny and Gross, 1974; Matthews and Gross, 1980; Geller *et al.*, 2012). The second hypothesis suggests that fires were activated by methane combustion originating from the tectonically active Dead Sea rift zone (Sokol *et al.*, 2010; Novikov *et al.*, 2013). Locally, melting of rocks of the protolith of the Hatrurim Complex can be observed, along with the formation of paralavas and slag-like rocks of different composition, forming bodies of different sizes from thin veins, ranging from gehlenite hornfels with a thickness of a few centimetres to large oval fields of diopside-bearing paralava with a diameter of



**Figure 2.** (a) Back-scattered electron (BSE) images of a sulfide nodule in which ferrodymolybdenite was discovered. The areas outlined are magnified in: Fig. 2b ('B') and Fig. 3a ('A'). (b) BSE image of nickelporphide and tetrataenite inclusions in troilite. The frame ('C') outlines the area magnified in Fig. 2c. (c) Reflected light image of the ferrodymolybdenite crystal has a inductive growth surface with troilite (arrow, simultaneous growth) and idiomorphism with respect to pentlandite and tetrataenite. Fdmol = ferrodymolybdenite; Krw = karwowskiite; Mol = molybdenite; Nic = nickelporphide; Pn = pentlandite; Tro = troilite; and Ttae = tetrataenite.

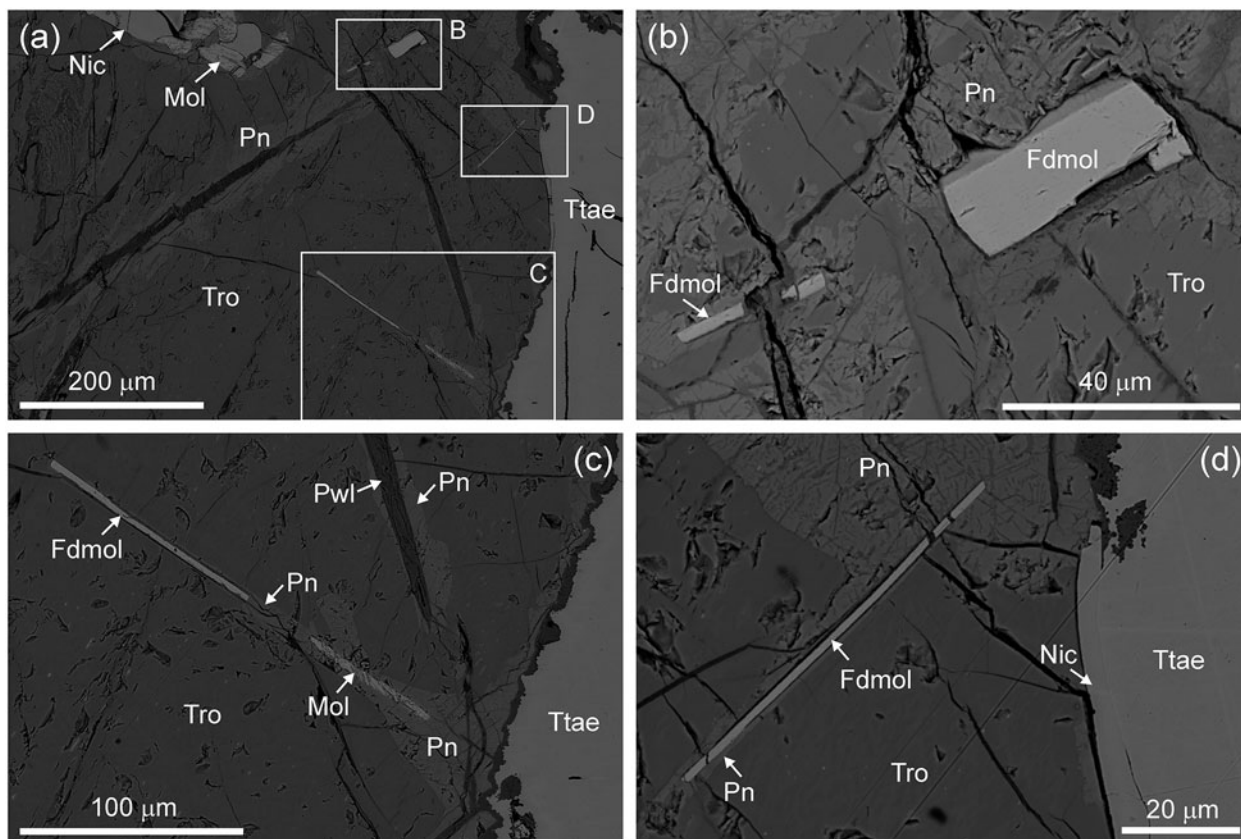
tens of metres. The near-surface character of the paralava generation processes determines the predominantly oxidised mineral composition of the paralavas, with minerals containing only  $\text{Fe}^{3+}$  (Galuskin *et al.*, 2017). In relatively rare cases, black reduced paralavas enriched with pyrrhotite are formed. The rarest are paralavas containing native iron and phosphides, as well as osbornite – an indicator of super-reduced conditions (Bittvín *et al.*, 2015; Galuskin *et al.*, 2022, 2023b, 2025; Futrzyński *et al.*, 2023). The hypothesis of phosphide and native iron formation as a result of reducing carbothermal reactions occurring at the boundary of hot paralava and thermally altered sedimentary rock enriched with graphitised and phosphorised organic residues has been proposed elsewhere (Galuskin *et al.*, 2022, 2023b, 2025). One of the rare examples of a reduced paralava is from Daba-Siwaqa, Jordan. Along with numerous aggregates of phosphides in the contact facies, this paralava features a sulfide nodule with inclusions of tetrataenite and nickelporphide, containing ferrodymolybdenite

crystals. It was found in the central part of the body (Galuskin *et al.*, 2025).

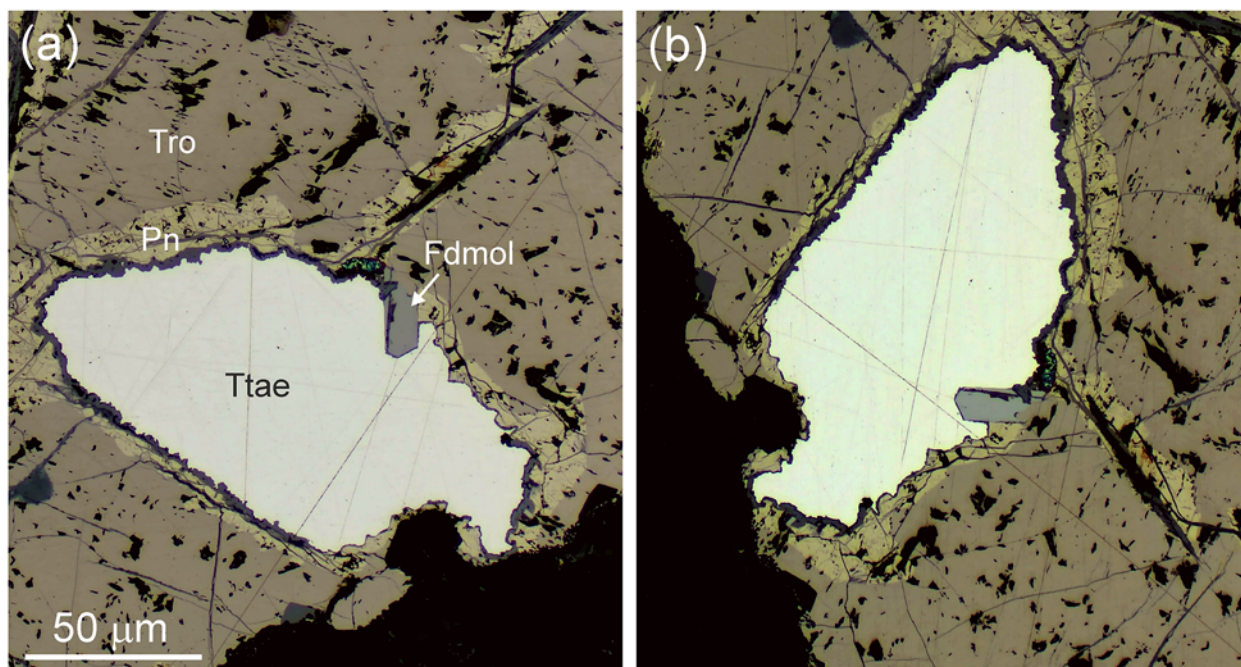
### Occurrence and composition

Ferrodymolybdenite was found in a small quarry (prospecting for phosphorite deposits) in the Daba-Siwaqa complex within the Transjordan Plateau, Jordan (31°22'01"N, 36°11'10"E), where a basalt-like paralava body ~30 metres in diameter is exposed in pyrometamorphically altered carbonate rocks of the Muwaqqar Chalk-Marl Formation. The paralava consists of diopside, anorthite, wollastonite, tridymite and small amounts of glass. Cristobalite is observed growing on tridymite crystals (Trd), exhibiting specific cracking patterns reminiscent of fish scales, indicative of the transition  $\text{Trd}_{\text{cub}} \rightarrow \text{Trd}_{\text{tet}}$  and a paralava generation temperature exceeding 1400°C (Galuskin *et al.*, 2023a). Fluorapatite, titanite and spinel of the





**Figure 3.** BSE images. (a) Fragment of sulfide nodule (from Fig. 2a) with ferrodymolybdenite crystals. The frames outline the areas magnified in Fig. 3b–d, respectively. (b) Relatively large ferrodymolybdenite crystal used for the single-crystal XRD study. (c, d) Fine lamellar crystals of ferrodymolybdenite (cross-section is  $\sim$  parallel to Z), usually with a thin film of pentlandite. Tro = troilite; Ttae = tetrataenite; Mol = molybdenite; Pn = pentlandite; Nic = nickelphosphide; Fdmol = ferrodymolybdenite; and Pwl = powellite.



**Figure 4.** Ferrodymolybdenite (centre right) in reflected light is (a) grey to light-grey with (b) a bluish tinge; the mineral exhibits strong birefringence. This crystal is also shown, magnified, in Fig. 2c. Tro = troilite; Ttae = tetrataenite; Pn = pentlandite; and Fdmol = ferrodymolybdenite.

**Table 1.** Reflectivity of ferrodimalybenite\*

| $R_{\max}$ | $R_{\min}$ | $\lambda$ (nm) | $R_{\max}$ | $R_{\min}$ | $\lambda$ (nm) |
|------------|------------|----------------|------------|------------|----------------|
| 35.7       | 34.2       | 400            | 36.9       | 35.1       | 560            |
| 35.8       | 34.3       | 420            | 37.1       | 35.7       | 580            |
| 35.9       | 34.3       | 440            | 37.3       | 35.9       | <b>589</b>     |
| 36.0       | 34.4       | 460            | 37.8       | 36.0       | 600            |
| 36.0       | 34.4       | <b>470</b>     | 38.2       | 36.2       | 620            |
| 36.1       | 34.4       | 480            | 38.9       | 36.8       | 640            |
| 36.2       | 34.4       | 500            | 39.0       | 37.0       | <b>650</b>     |
| 36.3       | 34.5       | 520            | 39.2       | 37.2       | 660            |
| 35.5       | 34.9       | 540            | 39.8       | 37.8       | 680            |
| 36.7       | 35.0       | <b>546</b>     | 40.0       | 38.1       | 700            |

\* The values required by the Commission on Ore Mineralogy (COM) are given in bold.

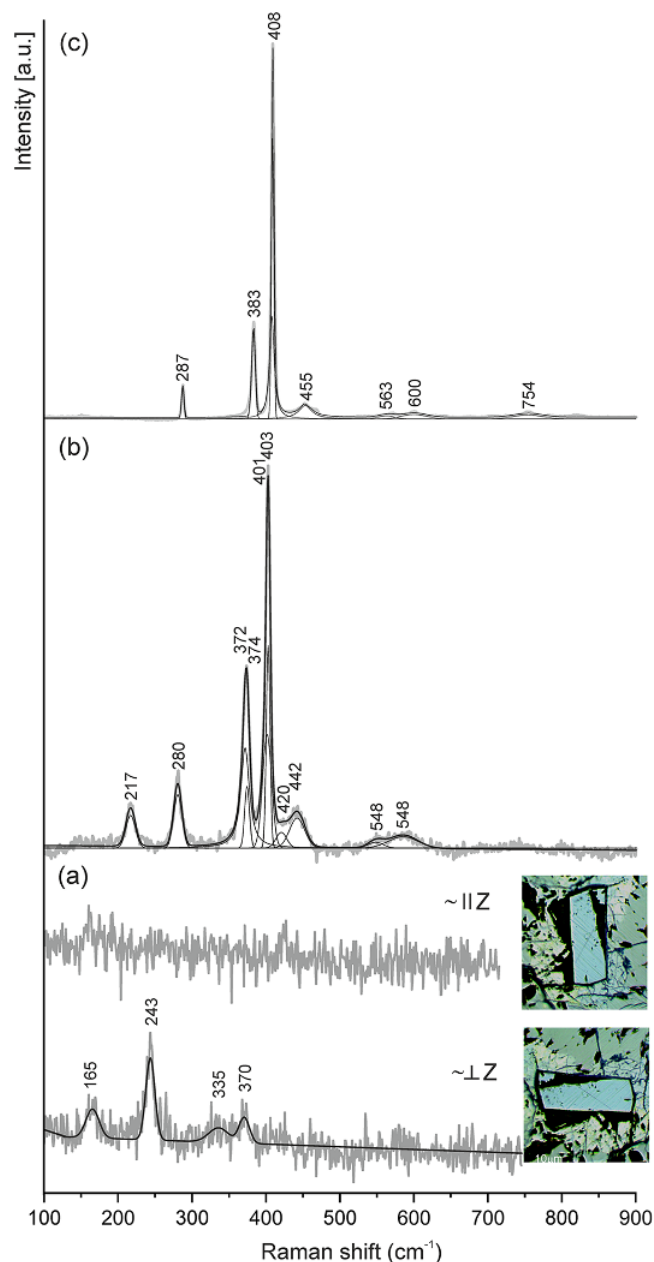
**Table 2.** Chemical composition (wt.%) of ferrodimalybenite

| Constituent | $n$   | S.D. | Range       |
|-------------|-------|------|-------------|
| Mo          | 49.30 | 0.39 | 48.96–49.95 |
| Cu          | 1.19  | 0.19 | 0.92–1.43   |
| Ni          | 0.58  | 0.14 | 0.39–0.77   |
| Fe          | 14.64 | 0.22 | 14.31–14.93 |
| S           | 33.90 | 0.14 | 33.69–34.03 |
| P           | 0.15  | 0.01 | 0.15–0.17   |
| Total       | 99.78 |      |             |

$n$  = number of spot analyses; S.D. = standard deviation.

magnesioferrite–magnetite–chromite series are the accessory minerals of the paralava. Ferrodimalybenite was identified in a sulfide nodule  $\sim 7$  mm in size (Fig. 1) found in the central part of the paralava body (Galuskin *et al.*, 2025). The sulfide nodule comprises troilite and pentlandite zones and gaseous bubbles (Figs 1b, 2a). Inclusions of tetrataenite, nickelphosphide, molybdenite, (rarely) galena and rudashevskyite are observed. The presence of crystals of the new mineral karwowskiite, with a composition of  $\text{Ca}_9\text{Mg}(\text{Fe}^{2+}_{0.5}\square_{0.5})(\text{PO}_4)_7$  (IMA2023–080), was detected on the boundary between the nodule and the paralava (Galuskin *et al.*, 2024). These crystals were absent in the groundmass of the rock. An analysis of the nodule minerals is presented in Galuskin *et al.* (2025). The composition of tetrataenite varies in the range  $\text{Fe}_{0.59}\text{Ni}_{0.39}\text{Cu}_{0.01}$  –  $\text{Ni}_{0.53}\text{Fe}_{0.44}\text{Cu}_{0.02}$  with a tendency for the Ni content to increase towards the edge of the grain. The composition of nickelphosphide also varies significantly, so the smallest crystals are characterised by the highest Ni content (up to 2.57 Ni per formula unit), while in the relatively large crystals in the centre the Ni content decreases to 2.06 Ni pfu, with a tendency to increase towards the edge of the crystal.

Ferrodimalybenite forms separate tabular crystals in a sulfide matrix, with dimensions ranging from 3  $\mu\text{m}$  to 20  $\mu\text{m}$  in the cross-section (Figs 2c, 3, 4). Ferrodimalybenite crystals are typically found in the troilite zone, with a thin pentlandite film commonly observed between ferrodimalybenite and troilite (Fig. 3b–d). Ferrodimalybenite is grey in colour with a dark grey streak. The mineral is opaque with a metallic lustre. Its Mohs hardness is  $\sim 3$ . Ferrodimalybenite crystals show cleavage in three directions: perfect on {001}, good on {100} and poor on {010}. Its tenacity is sectile, and its fracture is smooth. Its density,  $5.445 \text{ g}\cdot\text{cm}^{-3}$ , was calculated from the empirical formula and unit cell volume refined from the single-crystal XRD data. The mineral is slightly magnetic. In reflected light, it is grey to light grey with a bluish tinge. It is strongly anisotropic (Fig. 4). No internal reflections have



**Figure 5.** (a) Raman spectra of ferrodimalybenite; crystal orientation during the measurements is shown. (b) Raman spectrum of thermally altered ferrodimalybenite. (c) Raman spectrum of molybdenite from the sulfide nodule.

been observed. The reflectivity varies between 34.2% and 40.0% (Table 1).

The chemical composition of ferrodimalybenite is close to stoichiometric  $(\text{Fe}^{2+}_{0.99}\text{Cu}^{2+}_{0.07}\text{Ni}^{2+}_{0.04})_{\Sigma 1.10}\text{Mo}^{3+}_{1.94}(\text{S}^{2-}_{3.98}\text{P}^{3-}_{0.02})_{\Sigma 4.00}$  (Table 2). The simplified formula of ferrodimalybenite is  $(\text{Fe}^{2+}, \text{Cu}^{2+}, \text{Ni}^{2+})\text{Mo}_2\text{S}_4$ , leading to the ideal formula  $\text{FeMo}_2\text{S}_4$ . Interestingly, a 3  $\mu\text{m}$  grain of the Cu-analogue of ferrodimalybenite, with the composition  $(\text{Cu}_{0.78}\text{Fe}_{0.22})\text{Mo}_2\text{S}_{\sim 4}$  was found in the inclusion in the tetrataenite.

### Raman spectroscopy

In the Raman spectrum of ferrodimalybenite at laser beam polarisation ( $\sim \perp Z$ ,  $\sim$  parallel to molybdenite layers), a single weak

**Table 3.** Parameters for X-ray data collection and crystal-structure refinement for ferrodymolybdenite

| Crystal data               |                                                               | Refinement of structure                                   |                                         |
|----------------------------|---------------------------------------------------------------|-----------------------------------------------------------|-----------------------------------------|
| Formula from refinement    | Fe <sup>2+</sup> Mo <sup>3+</sup> <sub>2</sub> S <sub>4</sub> | Reflection measured                                       | 6695                                    |
| Crystal system             | Monoclinic                                                    | No. of unique reflections                                 | 2133                                    |
| Space group                | C2/c (no. 15)                                                 | No. of observed unique refl. [ <i>I</i> > 3σ( <i>I</i> )] | 1619                                    |
| Unit-cell dimensions       |                                                               | Refined parameters                                        | 64                                      |
| <i>a</i> (Å)               | 11.8249(8)                                                    | <i>R</i> <sub>int</sub>                                   | 0.0376                                  |
| <i>b</i> (Å)               | 6.5534(9)                                                     | <i>R</i> <sub>1</sub> / <i>R</i> <sub>all</sub>           | 0.0297/0.0469                           |
| <i>c</i> (Å)               | 13.0052(10)                                                   | <i>wR</i>                                                 | 0.0569                                  |
| β (°)                      | 114.474                                                       | Goof                                                      | 1.034                                   |
| <i>V</i> (Å <sup>3</sup> ) | 917.27(12)                                                    | Δρ <sub>min</sub> (e Å <sup>-3</sup> )                    | −1.563                                  |
| <i>Z</i>                   | 8                                                             | Δρ <sub>max</sub> (e Å <sup>-3</sup> )                    | 2.201                                   |
| Crystal size (mm)          | 0.040×0.030×0.010                                             |                                                           |                                         |
| Data collection            |                                                               |                                                           |                                         |
| Diffractometer             | SuperNova with Atlas CCD                                      | Weighting scheme                                          | $w = 1/[\sigma^2(F_o^2) + (0.0207P)^2]$ |
| Radiation wavelength       | MoKα, λ = 0.71073 Å                                           |                                                           |                                         |
| Min. & max. theta (°)      | 3.44, 36.16                                                   |                                                           |                                         |
| Reflection ranges          | −19 ≤ <i>h</i> ≤ 17                                           |                                                           |                                         |
|                            | −10 ≤ <i>k</i> ≤ 10                                           |                                                           |                                         |
|                            | −19 ≤ <i>l</i> ≤ 21                                           |                                                           |                                         |

**Table 4.** Atom coordinates (*x,y,z*), equivalent isotropic displacement parameters (*U*<sub>eq</sub>, Å<sup>2</sup>) for ferrodymolybdenite

| Site | Atom | <i>x</i>    | <i>y</i>    | <i>z</i>    | <i>U</i> <sub>eq</sub> |
|------|------|-------------|-------------|-------------|------------------------|
| Mo1  | Mo   | 0.09969(2)  | 0.39854(5)  | −0.00541(2) | 0.00574(8)             |
| Mo2  | Mo   | −0.09122(2) | 0.14863(5)  | 0.00741(2)  | 0.00592(8)             |
| Fe1  | Fe   | 0.25958(4)  | 0.37567(11) | 0.25116(4)  | 0.01127(15)            |
| S1   | S    | 0.03565(7)  | 0.38106(14) | 0.14654(7)  | 0.00712(16)            |
| S2   | S    | 0.24706(7)  | 0.37973(14) | −0.10464(7) | 0.00703(16)            |
| S3   | S    | 0.26566(7)  | 0.63228(14) | 0.11266(7)  | 0.00699(16)            |
| S4   | S    | −0.01610(7) | 0.12869(14) | −0.13516(7) | 0.00715(16)            |

band at 243 cm<sup>−1</sup> related to the stretching Mo–S vibration is observed (Fig. 5a). When laser beam power is increased, a dark spot appears on the surface of ferrodymolybdenite in which the Raman spectrum corresponds to that of molybdenite (Fig. 5b,c).

## Crystal structure

Single-crystal XRD and refinement showed the cation and anion sites are fully occupied. Details of data collection and structure refinement are given in Table 3, final atomic coordinates are summarized in Table 4, and anisotropic displacement parameters are given in Table 5. Selected bond lengths and bond valence sum (BVS) calculations are given in Table 6. The crystallographic information file has been deposited with the Principal Editor of

*Mineralogical Magazine* and is available as Supplementary material (see below).

Ferrodymolybdenite (C2/c; *a* = 11.8249(8) Å, *b* = 6.5534(3) Å, *c* = 13.0052(10) Å, β = 114.474(9)° and *V* = 917.27(12) Å<sup>3</sup>), like its synthetic analogue (C1c1; *a* = 11.8148(2) Å, *b* = 6.5499(1) Å, *c* = 13.014(2) Å, β = 114.455(1)° and *V* = 916.75 Å<sup>3</sup>) (Guillevis *et al.*, 1974; Vaqueiro *et al.*, 2002), crystallises with monoclinic symmetry and has a structure formed by layers of octahedra ∞(MoS<sub>2</sub>)<sup>−</sup> with columns of Fe<sup>2+</sup>-octahedra between them (Fig. 6a). Ferrodymolybdenite has a centrosymmetric structure, two types of Mo-octahedra and one type of Fe-octahedra (Fig. 6a,b). In the synthetic analogue there is no inversion centre and a reduction in symmetry is illustrated by the presence of four types of Mo-octahedra and two types of Fe-octahedra. The ferrodymolybdenite structure can be considered a molybdenite structure, where molybdenite layers are divided by parallel columns of Fe-octahedra. In natural molybdenite the ∞(MoS<sub>2</sub>)<sup>0</sup> layers consist of trigonal prisms (MoS<sub>6</sub>)<sup>8−</sup>, not octahedra. Distorted octahedra have only been found in synthetic molybdenite-1T Mo (Samy *et al.*, 2021). Mo-bearing sulfides with the general formula AMo<sub>2</sub>S<sub>4</sub> (*A* = V, Cr, Fe and Co) are isostructural with Cr<sub>3</sub>S<sub>4</sub> (space group C2/m). In this space group, the symmetry only allows distortions involving the formation of zigzag cation chains (Vaqueiro *et al.*, 2002). A reduction in symmetry from C2/m (archetype of the Cr<sub>3</sub>S<sub>4</sub> structure) to C1c1 for artificial FeMo<sub>2</sub>S<sub>4</sub> can be related both to its synthesis conditions (Canadell *et al.*, 1988) and to the deformation of its crystals during pulverisation for powder diffraction.

**Table 5.** Anisotropic displacement parameters (Å<sup>2</sup>) for ferrodymolybdenite

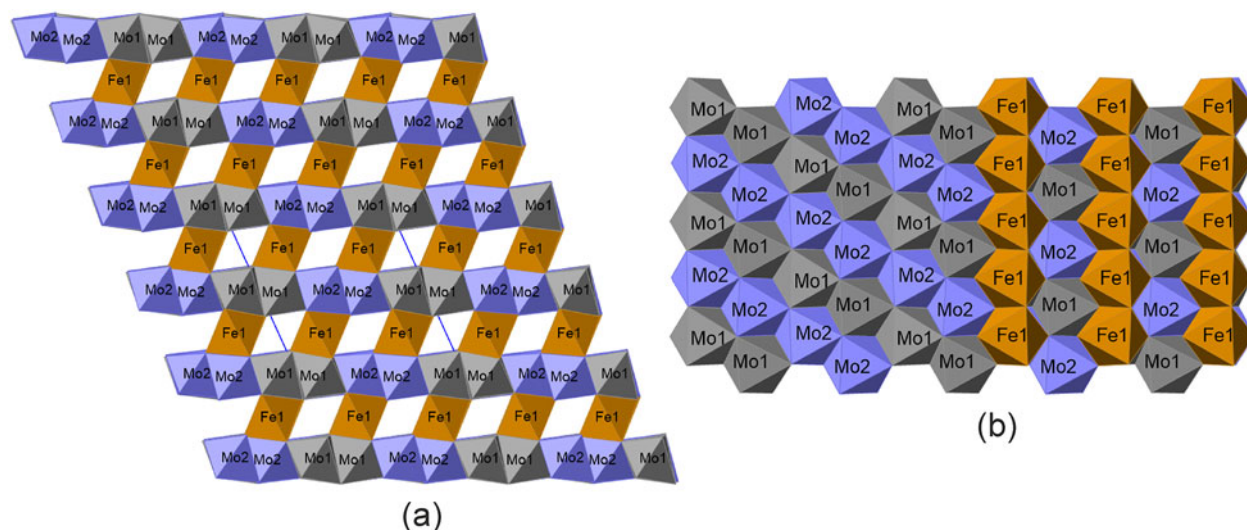
| Site | <i>U</i> <sup>11</sup> | <i>U</i> <sup>22</sup> | <i>U</i> <sup>33</sup> | <i>U</i> <sup>12</sup> | <i>U</i> <sup>13</sup> | <i>U</i> <sup>23</sup> |
|------|------------------------|------------------------|------------------------|------------------------|------------------------|------------------------|
| Mo1  | 0.00469(12)            | 0.00680(15)            | 0.00553(13)            | 0.00008(9)             | 0.00192(9)             | 0.00025(9)             |
| Mo2  | 0.00455(12)            | 0.00743(15)            | 0.00560(13)            | −0.00002(9)            | 0.00192(9)             | −0.00033(9)            |
| Fe1  | 0.0087(2)              | 0.0152(3)              | 0.0084(2)              | 0.00036(17)            | 0.00203(19)            | −0.00072(17)           |
| S1   | 0.0068(3)              | 0.0073(3)              | 0.0068(3)              | −0.0005(3)             | 0.0023(3)              | 0.0000(3)              |
| S2   | 0.0065(3)              | 0.0071(3)              | 0.0079(3)              | −0.0004(3)             | 0.0034(3)              | 0.0004(3)              |
| S3   | 0.0059(3)              | 0.0075(3)              | 0.0076(3)              | −0.0005(3)             | 0.0028(3)              | −0.0002(3)             |
| S4   | 0.0073(3)              | 0.0069(3)              | 0.0074(3)              | −0.0002(3)             | 0.0032(3)              | −0.0002(3)             |



**Table 6.** Selected bond distances (Å) and BVS (bond-valence sum) for ferrodymolybdenite

| Atoms   | distance  | atoms   | distance   | atoms   | distance   | Atom | BVS  |
|---------|-----------|---------|------------|---------|------------|------|------|
| Mo1–S1  | 2.3628(9) | Mo2–S1  | 2.3655(9)  | Fe1–S1  | 2.4247(10) | Mo1  | 2.87 |
| Mo1–S1  | 2.3941(9) | Mo2–S4  | 2.3682(9)  | Fe1–S4  | 2.4440(10) | Mo2  | 2.86 |
| Mo1–S4  | 2.4362(9) | Mo2–S4  | 2.4345(10) | Fe1–S3  | 2.4869(11) | Fe1  | 2.32 |
| Mo1–S3  | 2.4615(9) | Mo2–S3  | 2.4670(9)  | Fe1–S3  | 2.4915(10) | S1   | 2.12 |
| Mo1–S2  | 2.5513(9) | Mo2–S2  | 2.5646(9)  | Fe1–S2  | 2.5128(11) | S2   | 1.88 |
| Mo1–S2  | 2.5653(8) | Mo2–S3  | 2.5824(9)  | Fe1–S2  | 2.5176(10) | S3   | 2.08 |
| <Mo1–S> | 2.462     | <Mo2–S> | 2.464      | <Fe1–S> | 2.480      | S4   | 1.98 |

BVS calculated using *ECoN21* (Ilinca, 2022) and the bond-valence parameters of Brown and Altermatt (1985).



**Figure 6.** (a) The structure of ferrodymolybdenite ( $C2/c$ ) is built up by octahedral layers  $(Mo_2S_2)^{1-}$ , between which  $Fe^{2+}$  cations are located. Projection on (010). (b) Octahedral layers in ferrodymolybdenite are formed by two types of octahedra  $(MoO_6)^{9-}$ , between which there are columns of Fe-octahedra  $(FeO_6)^{10-}$  (only three columns are shown). Drawn using *CrystalMaker 2.7*® software.

Taking into account the systematic absences, it was found that two space groups are possible for ferrodymolybdenite, namely  $C1c1$  (no. 9) and  $C2/c$  (no.15). Statistical analysis of the intensities of the reflections using the *GRAL* programme implemented in the *CrysAlis<sup>Pro</sup>* package (Rigaku Oxford Diffraction, 2015) suggests that the space group should be centrosymmetric, i.e. the mean  $|E^2 - 1|$  value is equal to 1.202 (expected 0.968 centrosymmetric and 0.736 non-centrosymmetric), suggesting that the space group should be centrosymmetric (Howells *et al.*, 1950; Dolomanov *et al.*, 2009). The refinement of the crystal structures showed that in the space group  $C1c1$ , there were 127 necessary parameters and  $R_1 = 0.0332$ , whereas in the space group  $C2/c$ , there were 64 necessary parameters and  $R_1 = 0.0297$ . We believe that the studied crystal of ferrodymolybdenite, which formed at high temperature from sulfide melt at low pressure, has  $C2/c$  symmetry.

## Discussion

The melting of a heterogeneous protolith determined the formation of individual metal droplets in the paralava. The metal droplets were saturated with sulfur and phosphorus due to the flow of hot reduced gases, which are by-products of combustion processes (Galuskin *et al.*, 2025). A cristobalite fish-scale texture, which appears at the phase transition from cubic to tetragonal cristobalite has been found (Schmieder *et al.*, 2009). The cubic polymorph of cristobalite is stable at high temperature and low pressure and spontaneously transitions to the tetragonal form on cooling

(Damby *et al.*, 2014). The detection of the fish-scale texture suggests that the paralava had a temperature above 1400°C (Galuskin *et al.*, 2023a). The sulfide nodule minerals crystallised later than the rock-forming minerals of the paralava. The sulfide melt in the paralava migrated short distances through numerous gas channels in the incompletely crystallised paralava. We believe that karrowoskiite, the merrillite-group mineral found at the boundary of this sulfide nodule and the paralava, crystallised in the melt film on the amygdale wall before it was filled with sulfide melt (Galuskin *et al.*, 2024).

Nickelphosphide was one of the first minerals to crystallise in the sulfide melt, forming single crystals (Figs 1b, 2a). Ferrodymolybdenite was also one of the first to form, as it is usually represented by idiomorphic crystals in contact with sulfides or tetrataenite. In one case, an inductive growth surface was observed between a troilite and a ferrodymolybdenite crystal (Fig. 2c), which may indicate their synchronous growth for a certain time interval at a temperature of 1000–1100°C. The crystallisation of ferrodymolybdenite took place under reducing conditions in a monosulfide iron melt, which determined the stabilisation of  $Mo^{3+}$  in its structure. Molybdenite ( $Mo^{4+}$ ) is found in the pentlandite part of the nodule, which is a product of the crystallisation of iron-nickel sulfide melt (Figs 1b, 2a). Crystals of ferrodymolybdenite are located in the troilite part of the differentiated sulfide nodule. A thin layer of pentlandite is often noted between the ferrodymolybdenite crystals and pyrrhotite (Fig. 3c,d). Pentlandite has a non-stoichiometric composition of  $(Ni,Fe)_9S_8$ , suggesting the

presence of scattered electrons in its structure to compensate for the charge of the cations, and crystallisation from a melt/solution that is an electrolyte. The iron–nickel sulfide melt was the source of electrons that determined the appearance of a mineral with  $\text{Mo}^{3+}$ , which is unusual in natural systems:  $\text{Mo}^{4+} + e^- = \text{Mo}^{3+}$ .

**Supplementary material.** The supplementary material for this article can be found at [10.1180/mgm.2024.82](https://doi.org/10.1180/mgm.2024.82).

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**Competing interests.** The authors declare none.

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