The Hera orebody: A complex distal (Au–Zn–Pb–Ag–Cu) skarn in the Cobar Basin of central New South Wales, Australia

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Abstract
The Hera Au–Pb–Zn–Ag deposit in the southeastern Cobar Basin of central New South Wales preserves calc-silicate veins and remnant sandstone/carbonate-hosted skarn within a reduced anchizonal Siluro-Devonian turbidite sequence. The skarn orebody distribution is controlled by a long-lived, basin margin fault system, that has intersected a sedimentary horizon dominated by siliciclastic turbidite, with lesser gritstone and thick sandstone intervals, and rare carbonate-bearing stratigraphy. Foliation (S1) envelopes the orebody and is crosscut by a series of late-stage east-west and north-south trending faults. Skarn at Hera displays mineralogical zonation alongstrike, from southern spessartine–grossular–biotite–actinolite-rich associations, to central diopside-rich–zoisite–actinolite/tremolite–garnet-absent) association in remnant carbonate-bearing lithologies and sandstone horizons; the northern lodes also display zonation down dip to garnet present associations. High-T, prograde skarn assemblages rich in pyroxene and garnet are pervasively replaced by actinolite/tremolite–biotite-rich retrograde skarn which coincides with the main pulse of sulfide mineralization. The dominant sulfides are high-Fe–Mn sphalerite–galena–non-magnetic high-Fe pyrrhotite–chalcopyrite; pyrite, arsenopyrite; scheelite (low Mo) is locally abundant. The distribution of metals in part mimics the changing gangue mineralogy, with Au concentrated in the southern and lower northern lode systems and broadly inverse concentrations for Ag–Pb–Zn. Stable isotope data (O–H–S) from skarn amphiboles and associated sulfides are consistent with magmatic (or metamorphic) water and sulfur input during the retrograde skarn phase, while hydroxylates and sulfides from the wall rocks display comparatively elevated δ18O and mixed δ34S consistent with progressive mixing or dilution of original magmatic (or metamorphic) waters within the Hera deposit by unexchanged waters typical of low latitude (tropical) meteoritic waters. High precision titanite
(U–Pb) and biotite (Ar–Ar) geochronology reveals a manifold orebody commencing with high-T skarn and retrograde Pb–Zn-rich skarn formation at ≥403 Ma, Au–low-Fe sphalerite mineralization at 403.4 ± 1.1 Ma, foliation development or new mineralization at 390 ± 0.2 Ma followed by thrusting, orebody dismemberment at 384.8 ± 1.1 Ma and remobilization or new mineralization at 381.0 ± 2.2 Ma. The polymetallic nature of the Hera orebody is a result of multiple mineralization events during extension and compression and involving both magmatic and likely formational metal sources.

**Keywords**
Ar–Ar geochronology, Cobar Basin, mineral chemistry, skarn, stable isotopes, U–Pb geochronology

1 | INTRODUCTION

The Siluro-Devonian Cobar Basin is a major mining province in central New South Wales, Australia. The area hosts small to large polymetallic precious and base-metal deposits including those in the Cobar mineral field (Endeavor, CSA, New Cobar, Chesney, New Occidental, Peak and Perseverance mines) as well as the Hera mine and Nymagee deposit to the southeast. Past production and identified resources for the Cobar Basin exceeds 134.9 t Au, 1.91 Mt Cu, 3.46 Mt Zn, 1.8 Mt Pb and 3,832 t Ag (to the end 2015; Seccombe et al., 2017).

Orebodies of the Cobar Basin have been variously classified through time as reactivated/remobilized syn-depositional sub-exhalative to exhalative deposits (Suppel, 1984), epithermal and volcanogenic massive sulfide deposits (David, 2005, 2006), but more recently an orogenic sulfide–Au model has been applied, with mineralizing fluids derived from a mixing of basement and basinal fluids during burial (e.g., Lawrie and Hinman, 1998; Stegman, 2007; Seccombe et al., 2017; Downes and Poulson, 2018). Isotopic characteristics can provide a piece to the puzzle but do not give a unique solution, particularly in the absence of well constrained temporal and spatial discrimination of the results.

Limited work has been conducted on the early phase of mineralization for the orebodies in the main Cobar mining district due to the intense overprinting deformation fabrics, ore remobilization and overprinting phases of mineralization. Below we present a detailed petrographic, isotopic and geochronological study of the polymetallic Hera orebody in the southeastern Cobar Basin. Our data strengthens an intrusion-related, skarn genesis for the early phase of mineralization at Hera, as suggested by Fitzherbert et al. (2017a, b), and highlights the longevity and manifold nature of mineralization required to assemble a complex polymetallic Cobar Basin orebody.

2 | GEOLOGICAL SETTING

The Cobar Basin is an extensional basin that was deposited over basement of Ordovician turbidite and S-type Silurian granites during the late Silurian to Early Devonian (Figure 1). Deep-water parts of the basin interfinger with two volcanogenic troughs, to the south (Rast Group) and west (Mount Hope Group) and are flanked by limestone-bearing shelf sequences to the west (Winduck Group), as well as limestone and shallow water to subaerial syn-rift volcanic sequences in the east (Kopyje Group)
(Figure 1). The lower stratigraphic levels of the southeastern deep-water basin, shelf sequences and Ordovician basement sequences are intruded by syn-rift Early Devonian I-type plutons along the eastern margin of the basin and adjacent shelf sequences (Figure 1). The Cobar Basin is thought to have been inverted and deformed over 25 million years between 405 to 380 Ma (Perkins et al., 1994; Glen et al., 1996). Basin inversion was associated with reactivation of major basin/trough margin faults or near margin faults. Most Cobar Basin-hosted orebodies are concentrated in a belt along the eastern margin of the deep-water Basin (Figure 1). These deposits are mostly associated with zones of faulting, shear zone development and epizone hydrothermal alteration, although the

FIGURE 1  Group-level distribution of the Cobar Supergroup and underlying basement. Undercover interpretation is based on Fitzherbert et al. (2017a, b)
effects of deformation are relatively diminished to the south and hydrothermal metamorphic grades are significantly higher (hornblende hornfels facies).

Mineralization in the main (northern) Cobar mineral field is associated with greenschist facies high-strain zones and is linked with fluids that exploited major fault systems and regional lithological contacts. Correlative mineralization along these same fault systems and lithological contacts occurs on the southeastern margin of the basin, hosted within siliciclastic-dominant turbidite sequences of the lower Amphitheater Group and to a lesser extent shallow-water carbonate-bearing sequences of the Mouramba Group (Figure 1). The deposits are spatially associated with a major basin margin fault, the Rookery Fault, although the major orebodies are located immediately west of this fault system.

The genetic process(es) that have driven mineralization in the Cobar Basin remain unclear. Rifting and deposition of the mineralized sequences occurred after ca. 424 Ma (youngest age of exhumed basement rocks; Downes et al., 2016), but synchronous with intrabasin volcanism dated at 420–417 Ma (Downes et al., 2016). The youngest identified syn-rift intrusions/volcanics within, and immediately adjacent to the Cobar Basin are c. 411 Ma (Jones et al., 2020), although the upper parts of the volcanogenic Rast Trough, to the south, are dated at c. 412 Ma (Downes et al., 2016) and c. 409 Ma (Bull et al., 2008), signifying that rift-related volcanism extended into the Pragian (410.8 ± 2.8 to 407.6 ± 2.6 Ma—time scale of Gradstein et al. 2012). The youngest preserved basin fill is thought to be lowest Emsian c. 405 Ma (van der Wielen and Glen, 2008; Glen et al., 2016), potentially even Emsian (c. 400 Ma; Parrish et al., 2018) (Emsian, 407.6 ± 2.6 to 393.3 ± 1.2 Ma—time scale of Gradstein et al. 2012). Shallow-water sequences on the faulted eastern and western flanks of the Cobar Basin host Pragian to earliest Emsian limestone (sulcatus (lower Pragian) to dehiscenz (lowest Emsian) conodont zones (c. 406 Ma; Mathieson et al., 2016).

Inversion of the basin during the Tabberabberan Orogeny is thought to have been initiated c. 400 Ma (Glen et al., 1992), with continued compression until c. 380 Ma (Downes et al., 2016); a similar 20 million year interval for the Tabberabberan Orogeny is observed throughout south-eastern Australia (Fergusson, 2017). Fitzherbert et al. (2017a, b) demonstrated that sedimentary packages of the Cobar Basin did not undergo deep burial during this inversion and perceived biotite-zone greenschist facies and above regional metamorphic grades are related to, pre-foliation, hydrothermal highs associated with zones of mineralization. In fact, many Cobar Basin orebodies show evidence for mineralization and peak (highest temperature) hydrothermal alteration pre- or potentially early syn-foliation development, suggesting the initial mineralization event is ≥400 Ma. This does not preclude subsequent pulses of mineralization during periods of movement on large basinal faults with or without associated magmatism. Syn- to late-orogenic Pb–Zn–(Ag) mineralization has been recorded for several orebodies in the Cobar Basin, including the Endeavor, Manuka and even CSA. It is also well documented that the adjacent Cu–Au mineralization is significantly older (e.g., Lawrie and Hinman, 1998).

3 | DEPOSIT DESCRIPTION

The Hera orebody is an Au–Pb–Zn–Ag-rich orebody with a mineral resource (combined measured, indicated, inferred) totaling 2061 kt at 1.8 g/t Au, 2.8% Pb, 4.2% Zn and 34 g/t Ag (Aurelia Metals Ltd 2019 Annual Report to the Australian Stock Exchange, 30th October 2019 https://www.aureliametals.com/investors/annual-reports).

3.1 | Geometry

The Hera orebody is interpreted as a single fault-controlled, but grossly stratabound massive sulfide breccia body (McKinnon and Fitzherbert, 2017) with a general strike of 330°, dip of ~80° west and an overall plunge of 020° north at its base. A single satellite lens (Kershaws lode) occurs east of the main orebody. The orebody has been cut by a series of late-stage north-oriented faults striking between 005–015° and dipping 75° east, as well as by lesser east–west faults striking 110° and dipping 80° south. These late faults result in a series of steeply-dipping (80° west), west-stepping ore lodes that strike 337–344°, with apparent short strike lengths, apparent steep plunges and large vertical extent (Figure 2).

3.2 | Metal distribution

Hera has elevated Au, Ag, Cu, Pb and Zn. In terms of metal distribution and major sulfides, Au is relatively widespread at Hera, but there is a clear negative correlation between Au and Ag abundance with Ag elevated only in the upper North Pod and upper Far West lode (Figure 3). Galena and sphalerite are uniformly abundant. There is a negative correlation observed between Pb–Zn and Au abundance, and a positive correlation between Pb–Zn and Ag abundance (Figure 3). There is also a positive correlation between Ag–Pb–Zn abundance and the abundance of carbonate-hosted skarn, while Au has the inverse relationship (Figure 3).
Pyrrhotite and pyrite are concentrated in the upper parts of all lodes. Pyrrhotite is ubiquitous within the North Pod while pyrite is low in abundance except at the top of the Main South and Main North lodes (Figure 3). In addition, arsenopyrite is abundant in the North Pod while chalcopyrite is widespread toward the upper parts of most lodes and reports to a similar location as Fe-sulfides. The general sulfide paragenesis is presented in Figure 4 and is discussed further below.

3.3 Host rocks

The Hera orebody is hosted in the deep-water lower Amphitheatre Group of the Cobar Supergroup. The local geology around the mine sequences is dominated by thin bedded to laminated, fine-grained sandstone and siltstone. This fine-grained sedimentary facies is rich in quartz and albite, with abundant detrital white mica, biotite and Fe–Ti oxides. McKinnon and Fitzherbert (2017) described the destruction of host rock feldspar and detrital biotite (using hole rock chemistry and hyperspectral data) as far as 135 m away from the Hera orebody; detrital Fe–Ti oxides are also replaced by hydrothermal titanite over a similar distance from the orebody. Thick-bedded sandstone and gritstone intervals are prominent within the mineralized interval and although mineralogically similar to the fine-grained equivalents, much of the skarn alteration is focused in and around these beds. In addition, McKinnon and Fitzherbert (2017) described a
northerly increase in dolomitic zones within the orebody, with rare laminated dolomite clasts preserved within zones of massive sulfide within the North Pod.

### 3.4 Skarn and alteration mineralogy

Mineralization occurs in vein/breccia zones predominantly hosted by intensely silicified, siliciclastic sequences (siltstone, sandstone and gritstone). A halo of porphyroblastic hydrothermal biotite surrounds the orebody. This halo extends for 10s of meters into the host siliciclastic sedimentary rocks. The deposits high-T, prograde skarn mineralogy has two main textural forms, siliciclastic-hosted veins and breccia fill (siliciclastic-hosted skarn) hosted mainly within siltstone-rich sequences, and sandstone/carbonate replacement skarn (sandstone/carbonate skarn) hosted mostly within

**FIGURE 3** Long section of the Hera orebody looking west including metal distribution and general changes in gangue mineralogy. Pyrrhotite and pyrite vol% calculated according to McKinnon (2017)
sandstone, gritstone with rare carbonate (dolomitic) clasts. Prograde skarn is overprinted by two main pulses of retrograde alteration; a sulfide-rich phase with moderate-T hydrous-mineral-rich retrograde skarn and late-stage of low-T hydrous mineral-rich hydrothermal alteration. Figure 4 shows the general ore and gangue mineral paragenesis for the Hera orebody, while Figure 5 illustrates the spatial distribution of the main paragenetic phases and common crosscutting relationships.

Siliciclastic-hosted calc-silicate veins and breccia fill are, by far, the most commonly preserved form of high-T prograde skarn. In general, siliciclastic horizons and clasts are intensely silicified and enveloped by sulfide-rich breccias/veins. These frequently contain quartz-rich veins with garnet–zoisite–titanite–tremolite ± scheelite (low Mo) proximal to and within the sulfide orebody (Figure 6a). A biotite-rich mineralogy is also present distal to the orebody. Garnet within these veins displays strong chemical zonation (described below) from grossular-rich cores with calcite inclusions to spessartine-rich rims, which coincide with actinolite–zoisite and massive sulfide mineralization (Figure 6b). Veins are commonly enveloped by a 10–200 mm wide zoned alteration halo. This typically includes fine-grained garnet–titanite–anorthite–zoisite proximal to veins, then garnet–actinolite–anorthite–zoisite, followed by actinolite–biotite–anorthite and finally distal biotite–muscovite-rich zones that meld with the porphyroblastic biotite alteration halo surrounding the deposit. Garnet is absent from siliciclastic skarn in the North Pod.

Remnant zones of High-T, prograde skarn occur within and adjacent to the mineralized sequence and may occur as pods, as a replacement after sandstone horizons and/or after carbonate clasts/blocks (Figure 5). These comprise a grossular–quartz–diopside–actinolite–zoisite–anorthite ± carbonate (often dolomitic) assemblage (Figures 6c,d). The remnant high-T, prograde skarn zones display a broad mineralogical zonation along the entire strike of the orebody from a garnet-rich in the southern lenses, to

| Mineral species | Prograde skarn | Retrograde skarn | Deformation |
|-----------------|---------------|-----------------|-------------|
|                 | High-T        | Low-T           | Ductile     |
| **Ore minerals**|               |                 | Brittle     |
| Scheelite       |               |                 |             |
| Sphalerite      |               |                 |             |
| Galena          |               |                 |             |
| Arsenopyrite    |               |                 |             |
| Gold            |               |                 |             |
| Pyrhotite       |               |                 |             |
| Chalcopyrite    |               |                 |             |
| Pyrite          |               |                 |             |
| Ag/Sb salt/sulfide* |       |                 |             |
| **Gangue minerals** |           |                 |             |
| Garnet          |               |                 |             |
| Diopside        |               |                 |             |
| Wollastonite*   |               |                 |             |
| Anorthite       |               |                 |             |
| Zoisite/clinzoisite |           |                 |             |
| Biotite         |               |                 |             |
| Tremolite/actinolite |         |                 |             |
| Titanite        |               |                 |             |
| K-feldspar      |               |                 |             |
| Chlorite        |               |                 |             |
| White mica      |               |                 |             |
| Albite          |               |                 |             |
| Stilbite (Ca)/natrolite |     |                 |             |

Mineral*= Observed by Lay et al (2018)  
Mineral present  
Mineral remobilized  
Mineral dated

Figure 4: Combined ore and gangue mineral paragenesis for the Hera orebody.
pyroxene-rich assemblage in the Far West lenses (Figure 6e), and tremolite–biotite–anorthite ± (scheelite) assemblage (Figure 6f) in the upper North Pod while there is a garnet-bearing assemblage in the lower North Pod.

The retrograde skarn phase is ubiquitous, enveloping prograde skarn throughout the orebody. This hydrous retrograde skarn is sulfide-rich and shows a progression from moderate-T to low-T mineral associations. Moderate-T retrograde skarn is the dominant expression of this phase, often with a high-Fe sphalerite–galena–pyrrhotite ± chalcopyrite assemblage (Figures 4, 5, and 6g). Pyrite is locally abundant, particularly toward the upper margins of ore lenses with the gangue dominated by actinolite–tremolite–biotite±spessartine. Although in many parts of the North Pod this hydrous phase consists of monomineralic tremolite in addition to abundant sulfides. The sulfide mineralogy of North Pod is also more variable. Arsenopyrite is locally more abundant, in addition to the other sulfides, and there are silver sulfosalts and elemental silver in upper North Pod. Gold is often, but not exclusively associated with sulfides. It occurs as infill between breccia clasts of the hydrous retrograde skarn. Very localized K-feldspar–titanite-rich assemblages are developed in the altered host rock selvage to low-Fe sphalerite mineralization and appear to postdate the moderate-T retrograde skarn (Figure 6i).

The final stage of skarn-related alteration involves variable development of late-stage low-T, hydrous chlorite–muscovite-rich ± rare almandine assemblages, often associated with galena and low-Fe sphalerite.

The Hera orebody is enveloped by variably developed foliation with a biotite–tremolite–white mica–talc assemblage. Sulfide-rich stringers often parallel the foliation and may reflect a new phase of mineralization or, more likely, zones of remobilized ore (Figure 6j). Native gold can also be present in the biotite-rich foliation that envelopes the orebody. Biotite and tremolite within the foliation are variably retrogressed to chlorite-rich associations late in the ductile phase of deformation.

The orebody and overprinting foliation are cut by late-stage, west-verging reverse faults defined by quartz-rich veins with titanite, clinozoisite, chlorite, albite, adularia, tourmaline and zeolite (Figures 5 and 6k). These are generally barren but minor high-Fe sphalerite can be present, although a single fibrous tourmaline-rich vein was observed to contain abundant native gold. These
veins are predominantly chlorite-rich throughout the orebody but abruptly switch to being biotite-rich in the Far West Deeps and lower North Pod (below 535 level) zones.

Sulfide-rich zones that cross-cut foliation are commonly observed (Figure 6l). These may be a minor late mineralization stage or be zones of remobilized ore.

**FIGURE 6** (a) Garnet–quartz veins from siliciclastic skarn. Note the high-Fe sphalerite and galena above the vein. Sample from the Main North lode. (b) Photomicrograph of quartz–garnet–scheelite–zoisite vein. The inclusions in the core of garnet (red arrow) are calcite, while zoisite is the only inclusion phase to garnet rims and within the veins. (c) Silicified and garnet-flooded sandstone horizon. Note the late-stage quartz–chlorite-rich veins associated with partial replacement of skarn garnet. (d) Remnant garnet–diopside skarn block likely developed in a sandstone unit. Note the replacement (mostly of diopside) by retrograde tremolite and sulfide skarn around the margins of the high-T skarn fragment. Far West lode. (e) Pod of diopside-rich skarn from the Far West lode. Note the strong sulfide replacement around the margins. (f) Irregular biotite–scheelite veins within a siliclastic host from the North Pod. (g) Typical retrograde, sphalerite–galena–tremolite skarn. (h) Low-Fe sphalerite and gold forming the matrix to a brecciated tremolite-enriched hydrous skarn. The black spots are radiating clusters of galena that are typically seen replacing pyroxene within the hydrous skarn. Far West lode. (i) Margin of the low Fe-sphalerite–gold vein. Left is retrograde tremolite–galena skarn, right is high Fe-sphalerite–quartz–gold breccia matrix. The thin transition zone between the two is marked by replacement of the hydrous skarn by quartz–K-feldspar–low Fe-sphalerite–titanite. Far West lode. (j) Chalcopyrite within the biotite-rich S1 foliation (red dashed line) that envelopes the orebody. The foliation is well developed in the biotite-rich selvage around the orebody. Far West lode. (k) Late-stage quartz–clinozoisite (buff colored) titanite veins cross-cutting the S1 foliation that envelopes the orebody (red dashed line). (l) Massive galena mineralization cross cutting folded S1 foliation that envelopes the orebody (red dashed lines)

### 4  |  RESULTS

#### 4.1  |  Mineral chemistry

Microprobe analyses for garnet, amphibole, epidote, feldspar, pyroxene, biotite and chlorite were collected with a Cameca SX–100 Camebax microprobe at the Australian National University. Results are discussed below. Representative data are presented in Table 1.

**Garnet** is compositionally variable depending on textural location and may be complexly zoned. The andradite content of garnet is minimal with analyses normalizing to three components: $X_{grs}$ (grossular) = $\frac{Ca}{(Fe + Mn + Mg + Ca)}$, $X_{sps}$ (spessartine) = $\frac{Mn}{Fe + Mn + Mg + Ca}$ and $X_{alm}$ (almandine) = $\frac{Fe^{2+}}{(Fe^{2+} + Mn + Mg + Ca)}$. Garnet composition varies according to its spatial setting and timing (Figure 7a).

Garnet within prograde skarn associations, siliciclastic vein/breccia (quartz–low Mo scheelite–zoisite-rich; Figure 6a,b) and/or sandstone/carbonate replacement (garnet–pyroxene-rich skarn; Figure 6d,e) have an elevated grossular component compared to
**TABLE 1** Representative microprobe data represented as weight percent oxide and recalculated cation proportions

| Garnet | Garnet | Garnet | Garnet | Garnet |
|--------|--------|--------|--------|--------|
| **External to Main lode** | **External to Main lode** | **Within Main lode** | **Within Main lode** | **Within Main lode** |
| Core  | Rim   | Core  | Rim   | Core  | Shoulder | Rim |
| SiO₂   | 38.56 | 38.05 | 38.07 | 37.40 | 37.47 | 38.33 | 37.45 |
| TiO₂   | 0.38  | 0.33  | 0.36  | 0.19  | 0.08  | 0.30  | 0.17  |
| Al₂O₃  | 21.64 | 21.64 | 21.65 | 21.83 | 21.42 | 20.62 | 21.88 |
| Cr₂O₃  | 0.00  | 0.00  | 0.00  | 0.00  | 0.01  | −0.01 | 0.01  |
| FeO    | 4.86  | 5.94  | 6.11  | 10.92 | 6.06  | 6.77  | 11.31 |
| MnO    | 11.02 | 17.60 | 11.63 | 18.08 | 22.01 | 18.54 | 19.98 |
| MgO    | 0.07  | 0.18  | 0.05  | 0.43  | 0.23  | 0.69  | 0.53  |
| CaO    | 24.02 | 22.59 | 12.99 | 15.14 | 12.52 | 15.28 | 9.63  |
| Na₂O   | 0.00  | 0.00  | 0.00  | 0.00  | 0.00  | 0.00  | 0.00  |
| K₂O    | −0.01 | 0.00  | 0.01  | 0.01  | 0.00  | 0.00  | 0.00  |
| Total  | 100.56| 100.65| 100.47| 100.61| 100.29| 100.39| 100.96|
| Num ox | 24.00 | 24.00 | 24.00 | 24.00 | 24.00 | 24.00 | 24.00 |
| Si     | 5.96  | 5.96  | 5.91  | 5.93  | 5.95  | 6.04  | 5.94  |
| Ti     | 0.04  | 0.04  | 0.04  | 0.02  | 0.01  | 0.01  | 0.02  |
| Al     | 3.94  | 4.00  | 3.96  | 4.08  | 4.01  | 3.83  | 4.09  |
| Cr     | 0.00  | 0.00  | 0.00  | 0.00  | 0.00  | 0.00  | 0.00  |
| Fe²⁺   | 0.57  | 0.78  | 0.65  | 1.44  | 0.73  | 0.88  | 1.50  |
| Fe³⁺   | 0.06  | 0.00  | 0.14  | 0.01  | 0.07  | 0.01  | 0.00  |
| Mn     | 1.44  | 2.34  | 1.53  | 2.43  | 2.96  | 2.48  | 2.68  |
| Mg     | 0.02  | 0.04  | 0.01  | 0.10  | 0.06  | 0.16  | 0.12  |
| Ca     | 3.97  | 2.84  | 3.76  | 2.00  | 2.21  | 2.56  | 1.64  |
| Na     | 0.00  | 0.00  | 0.00  | 0.00  | 0.00  | 0.00  | 0.00  |
| K      | 0.00  | 0.00  | 0.00  | 0.00  | 0.00  | 0.00  | 0.00  |
| Total  | 16.00 | 16.00 | 16.00 | 16.00 | 16.00 | 16.00 | 16.00 |

| Biotite | Feldspar | Chlorite | Pyroxene | Epidote | Amphibole |
|---------|----------|----------|----------|---------|-----------|
| **Main South** | **North Pod** | **Within Main lode** | **Within Main lode** | **Unretrogressed skarn** | **Tremolite** | **Actinolite** |
| SiO₂    | 36.74    | 45.16    | 27.93    | 53.14   | 38.59     | 53.28     |
| TiO₂    | 1.25     | 0.01     | 0.02     | 0.00    | 0.01      | 0.01      |
| Al₂O₃   | 18.38    | 35.55    | 22.27    | 0.08    | 32.32     | 2.67      |

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|          | Biotite | Felspar | Chlorite | Pyroxene | Epidote | Amphibole |
|----------|---------|---------|----------|----------|---------|-----------|
|          | Main South | North Pod | Within Main lode | Within Main lode | Unretrogressed skarn within lode | Unretrogressed skarn within lode |
| Cr₂O₃    | 0.03  | 0.01 | 0.00 | 0.00 | 0.00 | 0.06 | −0.01 | −0.01 |
| FeO      | 16.51 | 8.42 | 0.05 | 0.12 | 20.63 | 20.94 | 7.92 | 7.33 |
| MnO      | 0.65  | 0.40 | 0.00 | 0.10 | 1.38 | 1.24 | 2.49 | 3.27 |
| MgO      | 12.62 | 21.84 | −0.04 | −0.03 | 17.80 | 17.49 | 12.52 | 12.46 |
| CaO      | 0.03  | 0.00 | 17.64 | 23.32 | 0.35 | 0.02 | 24.41 | 24.69 |
| Na₂O     | 0.08  | 0.04 | 1.22 | 0.50 | −0.01 | 0.01 | 0.00 | 0.03 |
| K₂O      | 8.77  | 8.18 | 0.01 | 0.02 | 0.01 | 0.17 | −0.01 | −0.01 |
| Total    | 95.05 | 94.75 | 99.60 | 99.15 | 90.37 | 91.21 | 100.54 | 101.10 |
| Num ox   | 24.00 | 24.00 | 8.00 | 8.00 | 24.00 | 24.00 | 6.00 | 6.00 |
| Si       | 6.01  | 6.67 | 2.08 | 1.93 | 2.81 | 2.83 | 1.99 | 1.96 |
| Ti       | 0.15  | 0.06 | 0.00 | 0.00 | 0.00 | 0.01 | 0.00 | 0.00 |
| Al       | 3.55  | 2.36 | 1.93 | 1.85 | 2.64 | 2.68 | 0.00 | 0.03 |
| Cr       | 0.00  | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 |
| Fe²⁺     | 2.26  | 1.10 | 0.00 | −0.32 | −4.13 | −4.15 | 0.23 | 0.18 |
| Fe³⁺     | 0.00  | 0.00 | 0.01 | 0.32 | 5.87 | 5.90 | 0.01 | 0.05 |
| Mn       | 0.09  | 0.05 | 0.00 | 0.00 | 0.12 | 0.11 | 0.08 | 0.10 |
| Mg       | 3.08  | 5.10 | 0.00 | 0.00 | 2.67 | 2.60 | 0.70 | 0.69 |
| Ca       | 0.00  | 0.00 | 0.87 | 1.16 | 0.04 | 0.00 | 0.98 | 0.99 |
| Na       | 0.03  | 0.01 | 0.11 | 0.05 | 0.00 | 0.00 | 0.00 | 0.00 |
| K        | 1.83  | 1.64 | 0.00 | 0.00 | 0.00 | 0.02 | 0.00 | 0.00 |
| Total    | 17.00 | 17.00 | 5.00 | 5.00 | 10.00 | 10.00 | 4.00 | 4.00 |
|          |         |         |         |         |         |         | 17.00 | 17.00 |

Note: Fe³⁺ recalculation is based on ideal stoichiometry.
almandine and spessartine that varies between $X_{\text{Alm}} = 8.29$–15.85, $X_{\text{SpS}} = 23.85$–41.36 and $X_{\text{Grs}} = 46.23$–65.76 (Figure 7a). Core to rim chemical zonation involves compositionally homogenous grossular-rich cores (sometimes with calcite inclusions) that display sharp boundaries with spessartine enriched rims, which also host zoisite and tremolite inclusions (Figure 6b). A thin outer-rim of elevated $X_{\text{Alm}}$, with decreased $X_{\text{Grs}}$ and increased $X_{\text{SpS}}$ values, may be present. These elevated $X_{\text{Alm}}$ outer rim analyses are grouped with the latest-stage low-T hydrothermal alteration. Where remnant garnet–pyroxene-bearing carbonate skarn is preserved, with limited hydrous retrogression, garnet tends to have a less well-developed chemical zonation and is dominated by grossular-rich compositions. Fine-grained garnet in alteration haloes adjacent to calc-silicate veins within siliciclastic sedimentary rocks are generally enriched in spessartine and almandine components and vary from $X_{\text{Alm}} = 6.31$–22.02, $X_{\text{SpS}} = 40.27$–64.60 and $X_{\text{Grs}} = 29.49$–49.94.

Garnet within mineralized retrograde skarn, quartz–tremolite-rich veins and carbonate–tremolite-rich sulfide breccia fill has an elevated spessartine component, varying from $X_{\text{Alm}} = 5.94$–18.69, $X_{\text{SpS}} = 41.76$–64.65 and $X_{\text{Grs}} = 23.85$–44.27. Core to rim chemical zonation involves a smooth decrease in the $X_{\text{SpS}}$ ratio and sympathetic increase in $X_{\text{Grs}}$ and $X_{\text{Alm}}$ values. A thin outer-rim of elevated $X_{\text{Alm}}$ with decreased $X_{\text{Grs}}$ and $X_{\text{SpS}}$ values may be present.

Fine grained garnet associated with late-stage low-T chlorite–quartz–sulfide veins is relatively enriched in almandine component varying from $X_{\text{Alm}} = 20.03$–28.57, $X_{\text{SpS}} = 33.27$–45.28 and $X_{\text{Grs}} = 23.85$–44.27. Garnet of this composition are generally small, granular and with no discernable chemical zonation and appears to correlate with almandine enriched garnet rims from all other settings.

**Amphibole** occurs in all stages of retrograde skarn and where it is developed in siliciclastic dominant parts of Hera (southern lodes) it varies from actinolite to manganoan-actinolite, Si p.f.u (per formula unit) ranges from 7.15 to 7.93 and XMg (Mg/Mg + Fe) ranges between 0.64 and 0.87 (Figure 7b). Manganese is elevated and ranges from 0.12 to 0.48 p.f.u. Lower Mn values tend to correlate with inclusions to garnet, while elevated Mn values tend to correlate with matted intergrowths of sulfide and amphibole external to garnet. Amphibole from the North Pod at Hera appears to have formed during both the prograde and retrograde skarn phases and is exclusively tremolite with Si p.f.u values ranging from 7.49 to 7.82, XMg values ranging from 0.89 to 1.0, and lower Mn values of 0.06–0.11 p.f.u. (Figure 7b). Amphibole associated with low-Fe sphalerite and nuggety gold in the Far West Lode is exclusively tremolite, with very low Mn values of 0.02–0.07 p.f.u.

**Epidote** is zoisite with $\text{Fe}^{3+}/(\text{Fe}^{3+} + \text{Al})$ p.f.u values of 0.00–0.1596. Epidote from remnant garnet–pyroxene skarn has values plotting at the low $\text{Fe}^{3+}$ end of this range. Allanite is visible optically, particularly in the North Pod, but was not analyzed as a part of this study. Late-stage, quartz vein hosted clinozoisite was not chemically analyzed as a part of this study.

**FIGURE 7** (a) Ternary plots of skarn garnet composition from Hera orebody. (b) Classification of amphiboles from Hera orebody based on Lee et al (1997). (c) Ternary plot of pyroxene composition from Hera orebody.
**Feldspar** associated with anhydrous skarn is anorthite to bytownite with \( \text{An (anorthite)} = \frac{\text{Ca}}{\text{Ca} + \text{Na} + \text{K}} \) values of 0.83–0.96. Orthoclase is present in association with low Fe-sphalerite and nuggety gold in the Far West lode. Albite (identified using XRD) is locally abundant in association with late stage bucky white quartz veins that crosscut the orebody.

**Pyroxene** from carbonate skarn is diopside (Figure 7c), although analyses display a manganese component with Mn p.f.u ranging from 0.07 to 0.10.

**Biotite** from outer zone of alteration haloes around siliciclastic-hosted skarn veins and in the porphyroblastic biotite zone surrounding the orebody has an XMg (Mg/(Mg + Fe)) ranging from 0.43 to 0.47. Biotite associated with tremolite/actinolite-rich retrograde skarn is more magnesian (XMg = 0.72). No biotite was analyzed from North Pod, although amphibole here trends to Mg-end member in the hydrous skarn from this zone and biotite is colorless to yellow brown in thin-section with low birefringence, consistent with the Mg-rich endmember. Biotite from the porphyroblastic halo around Hera orebody has a lower range in XMg values from 0.28 to 0.29.

**Chlorite** from late-stage hydrous alteration is predominantly ripidolite (ferroan clinochlore) with Si p.f.u values of 5.2–5.7 and \( 2\text{Fe}^{2+} + 3\text{Fe}^{3+} \) p.f.u values of 2.3–3.9. The lower values represent chlorite replacing inclusions in garnet.

### 4.2 | Stable isotopes

#### 4.2.1 | S-isotopes

Nineteen sulfur isotope analyses were carried out from retrograde Au-bearing skarn that immediately post-dates moderate-T retrograde skarn at Hera (seven analyses) and the moderate-T retrograde skarn in an adjacent mineralized zone (Kershaws zone;12 analyses) as part of the present study. These results are summarized in Table 2 and discussed below. The analytical procedure and tabulated results are included in the Supporting information.

S-isotope values from this study for Au-bearing low-Fe sphalerite-rich skarn at Hera average 5.4‰. Values for sphalerite range 5.1–6.0‰ (average 5.5‰ from six analyses) with a single value for galena–sphalerite of 4.6‰. The sulfur isotope values for the Kershaws zone are very similar and average 5.0‰. Pyrite from Kershaws has \( \delta^{34}S \) values between 5.1 and 5.4‰ (average 5.2‰ from four analyses), sphalerite has values between 5.0 and 5.5‰ (average 5.2‰ from three analyses), galena has values of 4.2 and 4.3‰ and there are
single $\delta^{34}S$ values for chalcopyrite of 5.4‰, for pyrrhotite of 5.5‰ and sphalerite–galena of 4.5‰. Sixty-one additional sulfur isotope analyses are also available from previous studies by David (2005), Mernagh (2008) and Downes and Poulson (2018). The combined dataset from all studies plot in a tight cluster (Figure 8) between 2.9 and 7.4‰ (mean 5.0‰). The combined dataset of S-isotopes is subdivided in terms of retrograde skarn (both moderate and low-T) and deformation related sulfides. Although there is some overlap and the variation is subtle, deformation related (foliation hosted and late stage cross cutting) sulfides tend to be 0.5–1‰ heavier than the equivalent retrograde skarn phase sulfides (Figure 8).

### 4.2.2 | O–H isotopes

Water $\delta^2$H–$\delta^{18}O$ values were determined using $\delta^2$H and $\delta^{18}O$ values for tremolite (from mineralized retrograde skarn), biotite, chlorite (both from the host rocks adjacent to skarn) and clinozoisite (from late cross-cutting veins) using estimated temperatures of last equilibration and adjusted for their mineral composition and are presented in Figure 9 (see Suzuoki and Epstein, 1976; Graham et al., 1980; Matthews et al., 1983; Graham et al., 1984; Graham et al., 1987; Fortier and Giletti, 1991; Vennemann and O’Neil, 1996; Cole and Ripley, 1999 and Saccocia et al., 2009). The estimation of water $\delta^2$H–$\delta^{18}O$ values from biotite can be problematical. Biotite–water exchange experiments of Suzuoki and Epstein (1976) show a systematic decrease in the hydrogen isotope mineral–water fractionation factor ($10^3\ln\alpha$) from 850 to 400°C. The experimental findings of Vennemann and O’Neil (1996) indicated a pronounced increase in $10^3\ln\alpha$ for temperatures between 400°C and 400°C. The inflection point between these two trends is around 350°C, although there is a mismatch in the compositions of biotite used in these experiments. We have used a uniform value of −38.1 for $10^3\ln\alpha$ to estimate water $\delta^2$H values for water–biotite pairs estimated to have exchanged in the range 300–400°C to accommodate this inflection point and to avoid extrapolating the results of Suzuoki and Epstein (1976) below their experimentally determined range. An XMg value of ~0.8 was determined for Mg-rich biotite from the skarn margin (from microprobe data), although biotite in the most magnesian assemblages, particularly from North Pod, may have values closer to XMg = 1. Extrapolation of the Suzuki and Epstein equilibria to 300–400°C to accommodate this inflection point and to avoid extrapolating the results of Suzuki and Epstein (1976) below their experimentally determined range. An XMg value of ~0.8 was determined for Mg-rich biotite from the skarn margin (from microprobe data), although biotite in the most magnesian assemblages, particularly from North Pod, may have values closer to XMg = 1. Extrapolation of the Suzuki and Epstein equilibria to 300–400°C to accommodate this inflection point and to avoid extrapolating the results of Suzuki and Epstein (1976) below their experimentally determined range. An XMg value of ~0.8 was determined for Mg-rich biotite from the skarn margin (from microprobe data), although biotite in the most magnesian assemblages, particularly from North Pod, may have values closer to XMg = 1. 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Extrapolation of the Suzuki and Epstein equilibria to 300–400°C to accommodate this inflection point and to avoid extrapolating the results of Suzuki and Epstein (1976) below their experimentally determined range.
4.3 U–Pb geochronology

Sensitive high-resolution ion microprobe (SHRIMP), isotope dilution thermal ionization mass spectrometry (ID-TIMS) and laser ablation inductively coupled plasma mass spectrometry (LA-ICP-MS) U–Pb dating was carried out on titanite from three different settings at Hera: (1) quartz–ilmenite–titanite vein that cross-cuts prograde skarn in the Far West lode (Sample JAF01), (2) remnant low-T retrograde, K-feldspar-altered skarn fragments associated with low-Fe sphalerite–titanite–Au-rich breccia (Sample JAF02) and, (3) thrust fault-related late-stage quartz–chlorite–titanite veins that crosscut the orebody (Samples JAF03 and JAF04). Grains of titanite 100 μm or greater in diameter (some broken from cm-scale grains) were liberated by hand-crushing. Titanite fragments from all samples are transparent and colorless, with occasional zones of fluid inclusions. Back-scattered electron images show that all fragments lack discernible zonation, reflecting the chemically homogeneous nature of the grains. Detailed analytical methods and results are outlined in the Supporting information.

Twenty-seven LA-ICP-MS U–Pb analyses were carried out on titanite grain fragments from sample JAF01. Measured U/Pb ratios were calibrated via measurements of the reference titanite MKED1 (Spandler et al., 2016).

FIGURE 10 Tera-Wasserburg plots showing U–Pb results for titanite samples and correlative photographs of actual dated samples from (a) Au-rich mineralization in the Far West lode in the Hera mine; (b) a quartz–ilmenite–titanite vein cross-cut garnet-rich skarn in the far west lode at Hera mine; (c and d) from hydrothermal quartz veins that partially dismember the Hera orebody. SHRIMP or LA-ICP-MS data is displayed in the main panel, and where collected ID-TIMS data in the inset. All data are plotted with 2 sigma uncertainties. Individual SHRIMP and LA-ICP-MS analyses are colored by U content. All ages are interpreted as the timing of titanite crystallization and/or closure of its U–Pb isotopic system.
The dataset is portrayed on a Tera-Wasserburg diagram (Figure 10a). The Pb-isotopic systematics for this sample suggests that the U–Pb system is undisturbed (within the precision of the analyses) and that the Devonian lower intercept represents the true age of titanite crystallization and/or U–Pb isotopic closure. This sample gave a date of 402.3 ± 3.6 Ma for the quartz–ilmenite–titanite vein that crosses high-T prograde skarn in the Far West lode (Table 3).

Between 25 and 37 SHRIMP U–Pb analyses were carried out on titanite fragments from sample JAF02, JAF03 and JAF 4 on the SHRIMP IIe instrument at Geoscience Australia, Canberra. All analyses were characterized by low U contents (2–107 ppm, median 35 ppm). Sample JAF03 has a bimodal distribution with some grains having very low U (7–11 ppm). Consequently, SHRIMP analysis for this sample targeted a subset of grains with higher U (57–93 ppm, median 75 ppm). By contrast, titanite from sample JAF04 had a wider range of U values (6–107 ppm, median 39 ppm). For both samples, Th/U is uniformly low (0.02–0.04, median 0.03 and 0.03–0.07, median 0.05 respectively). Sulfide-Au hosted skarn sample JAF02, is characterized by U values (2–69 ppm, median 27 ppm) that are mostly lower than for JAF03 and JAF04, but Th/U is markedly higher (0.15–1.25, median 0.80). Common 206Pb as a proportion of total 206Pb is high (8–73%, median 20%) for this sample as is usual, in Phanerozoic titanite, and displays a largely predictable relationship with U content. Measured titanite U/Pb ratios were calibrated via measurements of reference titanite MKED1 (Spandler et al., 2016) with no correction for common Pb. The results are portrayed on Tera-Wasserburg diagrams (Figure 10a–d) with each dataset defining a statistically coherent discordia array (p exceeds .05 in all three cases). In all three cases, the y-intercept of the discordia is indistinguishable from the range of Devonian bulk-Earth 207Pb/206Pb values as modeled by Stacey and Kramers (1975): that is, 0.864 at c. 419 Ma through 0.860 at c. 359 Ma, and direct Pb-isotope measurement on 11 galena samples from the Hera mine averaging 0.863 (Downes and Poulson, 2018).

This suggests that each of the U–Pb systems are undisturbed (within the precision of the analyses) and that each Devonian lower intercept represents the true age of titanite crystallization (and/or U–Pb isotopic closure). Sample JAF02 gave an age of 404.2 ± 5.1 Ma for the sulfide-rich skarn-hosted sample, while JAF03 and JAF04 gave dates of 384.3 ± 3.4 Ma and 382.5 ± 3.1 Ma (respectively) for the quartz-rich veins hosted by crosscutting, thrust related structures. These results are summarized in Table 3.

Titanite grains from samples JAF02, JAF03 and JAF04 were reanalyzed at the Boise State University Isotope Geology Laboratory using ID-TIMS, to obtain more precise U–Pb dates, and to evaluate the accuracy of the MKED1-based calibration of 206Pb/238U used in the SHRIMP dating. Five titanite fractions with similar chemical composition to those analyzed via SHRIMP were analyzed from each of the three samples. Common Pb corrections were made using measured 204Pb and the two-stage Pb isotope evolution model of Stacey and Kramers (1975) at the nominal sample age. In all three cases, the five analyses for each sample are indistinguishable within their analytical uncertainties and define weighted mean 206Pb/238U dates well within uncertainty of the lower-intercept dates determined from the SHRIMP-derived discordia arrays (Figure 10b–d; Table 3). Sample JAF02 gave an age of 403.4 ± 1.1 Ma, while JAF03 and JAF04 gave dates of 384.0 ± 0.6 Ma and 384.8 ± 1.1 Ma (respectively). These results are summarized in Table 3. The excellent agreement between all three pairs of results confirms the accuracy of the MKED1-based calibration of SHRIMP-determined 238U/206Pb in the unknowns. All ID-TIMS and SHRIMP results from statistically coherent age groupings and the SHRIMP and ID-TIMS data for each sample are within uncertainty of each other (Table 3).

**4.4 Ar–Ar geochronology**

Ar–Ar dating was carried out at the University of Melbourne noble gas laboratory, on two biotite separates...
from biotite-rich schist developed around the margins of the Hera orebody (samples NSW34-2 and NSW34-3). Detailed analytical methods and results are outlined in the Supporting information. Age spectra for both separates are presented in Figure 11 and results are summarized in Table 4. The spectra are characterized by anomalously young apparent ages for the initial heating step (339.1 ± 1.2 Ma for NSW34-2, 346.0 ± 0.6 Ma for NSW34-3). Followed by a gradual increase from ~385 to ~389 Ma for low to intermediate steps (385.62 ± 0.45 Ma to 388.87 ± 0.41 Ma for NSW34-2; 385.48 ± 0.33 to 388.41 ± 0.37 Ma, for NSW34-3). The mid- to high-temperature steps yield identical plateau ages of 390.03 ± 0.23 Ma (59.5% 39Ar release for NSW34-2) and 389.60 ± 0.23 Ma (57.2% 39Ar release, NSW34-3).

The anomalously low age yielded by the first heating step of each aliquot is most likely an artefact of argon recoil. The “staircase” portion of the spectra (385–389 Ma), may reflect either partial resetting at or after 385 Ma by a later thermal event, and/or be an artefact of argon recoil. Regardless, the mid- to high-temperature heating steps yield extremely concordant results (both within and between aliquots), and as such the plateau ages are considered to reflect the cooling age of the biotite or the crystallization age if crystallization occurred ≤300°C (i.e., below biotite 40Ar/39Ar closure temperature; McDougall and Harrison, 1999).

Downes and Phillips (2018) reported 40Ar/39Ar step-heating age spectra for undeformed muscovite laths and rosettes intergrown with sulfides (galena–pyrrhotite–pyrite) at Hera (drill hole HRD032; 514 m). The muscovite showed a well-defined plateau age of 381.9 ± 2.2 Ma. Table 4 summarizes the 40Ar/39Ar results from the two studies.

5 | DISCUSSION

5.1 | Hera mineral chemistry, stable isotopes, and fluid evolution: A magmatic-related distal skarn?

Distinctive petrographic and mineral chemical features of the Hera orebody (e.g., sub-calcic garnet, pronounced Mn and prevalence of Fe2+ in skarn minerals, as well as low Mo in scheelite) in combination with the absence of a thermal aureole are consistent with primary mineralization at Hera being related to the retrograde phase of a reduced (pyrrhotite-stable) distal Zn-skarn system as described by Einaudi et al. (1981) and Meinert (1992).

Prograde and retrograde skarn mineral assemblages vary from Ca–Mg enriched in the north (diopside, tremolite, phlogopite, talc, anorthite), to comparably Ca–Fe enriched in the south (garnet, actinolite, biotite). This change in mineralogy in part reflects changing host lithology and ultimately changing bulk chemistry of the reactive sequence, with Mg enriched carbonate-bearing sequences most prevalent in the north and exclusively siliciclastic, comparably Fe-enriched (siltstone-rich) sequences in the south. Another factor in the complex mineralogy and mineral chemistry of the Hera distal skarn is the inferred low XCO2 and potential decrease in XCO2 during skarn evolution. Carbonate preservation is poor, and it has been mostly consumed to form calc-silicate minerals during skarn formation. Figure 12 illustrates the T–XCO2 stability region for both prograde and retrograde mineral associations most applicable to the Mg-enriched calc-silicate assemblages of the northerly lodes (Far West and North Pod) of the Hera skarn.

| Sample       | Age (Ma, ±95% CI) | MSWD | p     | % 39Ar release | Mineral   |
|--------------|------------------|------|-------|----------------|-----------|
| HRD018       | 381.9 ± 2.2      | 1.4  | .24   | 65.5           | Muscovite |
| NSW34-2      | 390.03 ± 0.23    | 0.87 | .48   | 59.5           | Biotite   |
| NSW34-3      | 389.60 ± 0.23    | 1.6  | .20   | 57.2           | Biotite   |
Changing mineral parageneses imply the garnet–pyroxene ± wollastonite (see Figure 4) bearing prograde skarn phase of the Far West lode initially developed at XCO₂ ≥ 0.05 in the field of calcite stability (preserved mostly as inclusions in garnet), but XCO₂ rapidly decreased into the field of zoisite stability in the presence of garnet and diopside. Finally, decreasing temperature from south to north along the length of the Hera orebody (and potentially up plunge in the North Pod) can account for the loss of garnet and pyroxene and prevalence of amphibole in the upper North Pod. Peak prograde skarn temperature potentially varied from 400 to 430°C for the upper North Pod and 430 to 500°C for the Far West lode (Figure 12). Decreasing temperature and mineralizing fluid ingress during the retrograde skarn phase also resulted in hydrous, amphibolite-rich associations and the replacement of peak garnet–pyroxene-rich associations in Far West lode during retrograde skarn development (Figure 12). Temperature ultimately decreased to <350°C where chlorite-rich epizonal mineral assemblages formed.

The sulfur isotopic characteristics of the Hera orebody are unusual in the context of the Cobar Basin. Variation in δ³⁴S is narrow and low (2.9–7.4‰) consistent with δ³⁴S for granitic rocks (Hoefs, 2004), while most other Cobar Basin orebodies vary between 5 and 12‰ (some up to 20‰). Whole rock δ³⁴S for selected Devonian intrusive and volcanic rocks from the Cobar Basin range between 4 and 6.2‰ (Downes and Poulson, 2018), while reduced seawater sulfate associated with the Cobar Basin should be 10–15‰ (Seccombe et al., 2017). The narrow range and low S-isotope values, for Hera are most consistent with a Devonian intermediate to felsic igneous reservoir and do not support the interpretation that significant sulfur was sourced from a fractionated Early Devonian seawater sulfate reservoir. Interestingly, when sulfur isotopes for Hera are divided into those associated with skarn and those associated with overprinting deformation, δ³⁴S for the latter is consistently 0.5–1‰ higher. This increase in δ³⁴S may reflect input of heavier sulfur form the Cobar Basin sedimentary rocks during deformation and basin inversion.

Calculated water δD–δ¹⁸O values for retrograde skarn minerals (tremolite) from within the massive sulfide breccia at Hera form a tight array, being typical of magmatic or metamorphic water (Figure 9). There is trend toward increasing δD values, with minor shift in δ¹⁸O values for lower temperature minerals developed in the altered, siltstone-rich margin to the orebody (i.e., biotite) and/or in later paragenetic stages (chlorite) (Figure 9). Calculated δD–δ¹⁸O values for some retrograde skarn phase biotite in the wall rocks to the Hera orebody, as well as overprinting deformation related biotite and chlorite defining the foliation around the orebody fall outside the field of magmatic and metamorphic waters from Taylor (1997) and require an external source of unexchanged waters to account for the observed trend. It is likely that original magmatic or metamorphic water of the retrograde skarn phase underwent progressive mixing or dilution late in the skarn forming event and more likely during the early stages of deformation by unexchanged waters with δD–δ¹⁸O values typical of low latitude (tropical) meteoritic waters around the margins of the massive sulfide breccia—this is consistent with the paleogeographic setting of the Cobar Basin during the late Silurian–Early Devonian (see Domeier and Torsvik, 2014).

The features of skarn mineralization, including the preservation of prograde, high-T skarn assemblages hosted within sub-greenschist facies basin sedimentary rocks (Fitzherbert et al., 2017a, b) and thermal changes reflected by changing skarn mineralogy (south to north and down plunge) at the Hera orebody are inconsistent with the current model for formation of Cobar-type deposits through deep burial and regional-scale ingress of basement and basin derived fluids during basin inversion (Stegman, 2001). Although no direct link to magmatic
rocks has been established at Hera, the features listed below are suggestive of magmatic heat and fluid/metal/sulfur input(s) to the Hera mineral system:

- Rapid changes in skarn mineralogy and the large thermal contrast between distal basin and skarn mineralization reflecting proximity to major flow channels, possibly reflected by the pyroxene-rich Far West Lode (see Einaudi et al., 1981; Meinert, 1992; Fitzherbert et al., 2017a, b).

- $\delta^{34}S$ values form a very tight cluster between 2.9 and 7.4‰ (average 5.0‰) consistent with Devonian magmatic sulfur from the Cobar region.

- $\delta^D–\delta^{18}O$ values are consistent with progressive mixing or dilution of original magmatic/metamorphic waters within the Hera deposit by unexchanged waters typical of low latitude meteoritic waters.

- Pb–model ages of 412–397 Ma for galena from the Hera orebody (Downes and Poulson, 2018) agree well with U–Pb age of 403.4 ± 1.1 for hydrothermal titanite from Zn–Au-skarn at Hera (i.e., Pb-source age and the age of mineralization are the same). Such a scenario can be most simply explained by the presence of a ≥ 403 Ma distal magmatic influence.

- Fitzherbert et al. (2017a, b) estimated sub-greenschist facies burial metamorphic conditions of $T = 200–250^\circ C$ for unaltered host rocks at Hera, with an increase to peak temperatures of ≥400°C (potentially ≤500°C) for the anhydrous prograde skarn associated with the orebody. This 150–200°C (potentially 250–300°C) thermal contrast between host rocks and skarn is inconsistent with metamorphic-reaction skarn, and more consistent with a distal igneous-metasomatic skarn (see Einaudi et al., 1981).

5.2 Timing of mineralization within a regional tectonic framework

High precision geochronology suggests the Hera orebody has had a protracted history of mineralization/remobilization from initial fault controlled distal base-metal skarn at ≥403 Ma, to Zn–Au-rich mineralization at 403 Ma, inversion initiation and ductile foliation development and ore remobilization/new mineralization at c. 390 Ma, brittle reverse faulting with orebody dismemberment and late inversion-related remobilization/mineralization at 381 Ma. The Cobar Basin has the highest metal endowment of the Siluro-Devonian basins in New South Wales (see Greenfield et al., 2015; updated figures for Cobar Basin, Seccombe et al., 2017). Perhaps a key to this endowment is the plurality of mineralization along the major eastern fault corridor, Rookery Fault. Rather than relying on a single ore forming event it is likely that the Hera deposit developed through successive mineralizing, remobilization/concentration events over ~20 Ma.

A generalized model for the opening of the Cobar Basin involves a rift phase from 420 to 415 Ma followed by a sag phase between 415 and 406 Ma (van der Wielen and Glen, 2008, Glen et al., 2016) and possibly as late as
400 Ma (Parrish et al., 2018). A recent review of conodont fauna in limestone of the Cobar Basin by Mathieson et al. (2016) places allochthonous limestone blocks hosted within a rift to sag phase (Biddabirra Formation; Figure 1) transitional facies in the Pragian at c. 410–408 Ma. An estimated 6–8 km of sag phase turbidite overlies the Biddabirra Formation (Glen, 1987). Deposition in the flanking shallow-water sequences (Winduck Shelf in the west and Kopyje Shelf [Yarra Yarra Creek Group] in the east; Figure 1) continued into the Emsian c. 406 Ma (Mathieson et al., 2016). Bull et al. (2008) and Downes et al. (2016) dated rift-related, deep-water A-type volcanics (Rast Trough; Figure 1) at c. 412–409 Ma (Figure 1). The Hera skarn may have formed due to fault movements during the later stages of sag phase of the basin deposition, but magmatic rocks this young are so far undiscovered in the Cobar Basin (see Figure 13 for an interpreted cross section through the ca. 410–400 Ma late sag phase Cobar Basin).

The Rookery Fault is a splay off a crustal-scale master fault system, the Gilmore Fault (Glen, 1982). The Gilmore Fault separates two tectonic entities in New South Wales; the central Lachlan Orogen (host of the Cobar Basin) and the eastern Lachlan Orogen. Basins (Jemalong Trough; Figure 16) preserved in the eastern Lachlan Orogen directly adjacent to the Gilmore Fault record a protracted history of rifting and bimodal magmatic activity through the Pragian to the middle Emsian c. 410–400 Ma (Lyons et al., 2000; Raymond et al., 2000, 2000a). Further east (Yass Shelf, Hill End Trough, Goulburn Trough and Bungonia Shelf; Figure 14) voluminous I-type silicic magmatism continued well into the early Emsian and was followed by a period of quiescence and limestone accumulation until c. 400 Ma (e.g., Thomas and Pogson, 2012). We propose that far-field forces associated with Pragian and potentially early Emsian tectonism in the eastern Lachlan Orogen have been translated as movement along the Gilmore and linked Rookery Fault systems. Distal skarn developed within the Cobar Basin, likely associated with limited magmatism that was focused along this crustal-scale fault system. Fault-focused magmatism has occurred during prior movements on the Gilmore and Rookery Faults, including the Avenall Metabasic Complex (415–411 Ma; Cairns, 1997; Kositcin, 1999) that stitches the Gilmore Fault to the south of Cobar, as well as fault localized dykes/plutons (415–410 Ma; Downes et al., 2016; Jones et al., 2020) within the Rookery Fault system. It is credible that extension reflected by continued rifting and magmatism in the east Lachlan Orogen from 415 to 400 Ma translated into periodic movements and fault focused diking/plutonism on the
predominantly strike-slip Gilmore-Rookery Fault system.

Inversion of the Cobar Basin is thought to have occurred between 400 and 380 Ma (Glen et al., 1992). Our dating suggests that basin inversion progressed from ductile deformation and foliation generation c. 390 Ma to brittle thrusting at c. 384 Ma. There is certainly evidence of sulfide and Au deposition post-dating the foliation at Hera but based on the above evidence it is likely the Ar–Ar white mica age of 381.9 ± 2.2 Ma for mineralization (Downes and Poulson, 2018) represents a cooling age toward the end of the Tabberabberan Orogeny inversion. In saying this, Pb–Zn mineralization is clearly associated with the Tabberabberan Orogeny and movement of basinal fluids within other orebodies of the Cobar Basin (e.g., Endeavor mine; Lawrie and Hinman, 1998; Sun et al., 2000). The late-stage Pb–Zn mineralization at Hera may potentially be of this generation.

5.3 Recognition of a distal skarn district

The link between faults, stratigraphy and mineralization at Cobar has been discussed before (e.g., Berthelsen, 2006), but the relatively recent development of the Far West lode and the North Pod at Hera has uncovered a previously unknown skarn altered, sandstone/gritstone/carbonate-bearing stratigraphy. In light of the newly described skarn association at Hera, reconnaissance studies have revealed that skarn mineralization in the southeastern Cobar Basin is relatively wide-spread (Fitzherbert et al., 2017a, b). Garnet–anorthite-rich skarn alteration is associated with the Cu-rich Nymagee deposit, 5 km north of Hera (Fitzherbert et al., 2017a, b) and garnet-rich calc-silicate veins are present at the Happy Jacks prospect a further ~2.5 km north (along strike) from Nymagee (Fitzherbert et al., 2017a, b). Mineralization and garnet-rich skarn associations are also recorded to the south of Hera, at Hebe prospect (Fitzherbert et al., 2017a, b). This distribution of skarn and skarn veins is associated with the same fault system and at the same stratigraphic level, being controlled by a single (>7.5 km strike length) horizon adjacent to a major basin margin fault. Allochthonous carbonate blocks are also present along the western margin of the basin. These include the Norma Vale prospect, where pyroxene-rich skarn is associated with Cu mineralization at the base of a large allochthonous limestone block within the Shume Formation (Figure 1; Fitzherbert et al., 2017a, b). The Shume Formation and correlative Biddabirra Formation both host allochthonous limestone and are stratigraphically higher than the Hera–Nymagee allochthonous horizon suggesting there are a number of prospective allochthonous rock packages within the deep-water Cobar Basin stratigraphy. The lower Cobar Basin sequences comprise shallow water reef to shelf sequences developed at the early-stage of rifting. These lower basin sequences host abundant conglomerate and carbonate, and with a shift in paradigm to a skarn mineral system model these shelf sequences become highly prospective for skarn mineralization.

6 CONCLUSIONS

The Hera orebody preserves primary distal skarn mineralization and has petrographic and mineral chemical characteristics consistent with a reduced, polymetallic siliciclastic/calcic skarn. Skarn mineralization in the Cobar Basin occurs where long-lived basin margin fault systems intersected stratigraphy that contains coarse-grained sandstone/gritstone horizons and rare carbonate clasts. The O-H isotopic characteristics of the Hera orebody are consistent with magmatic/metamorphic water that mixed with cooler low latitude basinal fluids, while sulfur isotopic characteristics imply mixed magmatic/basin sulfur input. The coincidence of the Pb-model ages for Hera with U–Pb titanite ages for mineralization implies the early Emsian ≥405–402 Ma was a period of movement along the Rookery Fault. This movement, in conjunction with fault focused magmatism likely gave rise to structurally hosted distal skarn at Hera. It is plausible that multiple movement events and potentially varying magmatic/basin metal input has resulted in the complex, polymetallic nature of the Hera skarn. A ductile foliation associated with basin inversion is dated at c. 390 Ma and the Hera orebody was dismembered by brittle faulting at c. 384 Ma. Late-stage Pb–Zn mineralization that cross-cuts the foliation is present at Hera, has been dated at c. 381 Ma and may reflect further basin derived mineralization during post-orogenic relaxation. Skarn mineralogy has been described at several prospects and deposits up to 7 km away from the Hera orebody implying that a larger and highly prospective distal skarn system occurs in the region.

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The data that supports the findings of this study are available in the supplementary material of this article.

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**SUPPORTING INFORMATION**

Additional supporting information may be found online in the Supporting Information section at the end of this article.

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