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From source to surface: clues from garnet-bearing Carboniferous silicic volcanic rocks, Iberian Pyrite Belt, Portugal

A. Cravinho¹ · D. Rosa² · J. M. R. S. Relvas¹ · A. R. Solá³ · I. Pereira^{4,5} · J.-L. Paquette⁵ · M. L. Borba⁶ · C. C. G. Tassinari⁶ · D. Chew⁷ · F. Drakou⁷ · K. Breiter⁸ · V. Araujo⁹

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Abstract

This work investigates the relationships between partial melting, melt extraction, pluton growth and silicic volcanism in garnet-bearing felsic volcanic rocks that were extruded in the Iberian Pyrite Belt, at ca. 345 Ma. The garnets are of peritectic origin, displaying textural and chemical features of disequilibrium crystallization during partial melting reactions involving biotite at high temperatures (up to 870 °C) in the middle-lower crust. Major element composition suggests compositional equilibrium with the entrained and pinitized peritectic cordierite, but reveals some subsequent homogenization by diffusion. Trace element maps and spot analyses of garnet show, nonetheless, significant trace element variations, reflecting biotite and Y-REE-P-rich accessory phase breakdown during partial melting reactions. Peritectic garnet and cordierite growth resulted in the preservation of Th- and Y-rich prograde suprasolidus monazite, which constrains the timing of partial melting of the metapelitic protolith at *ca.* 356.8 ± 2.4 Ma. The zircon cargo further shows that a significant amount of zircon crystals from previously crystallized felsic melts were also remobilized and erupted. These were likely stored in an upper crustal pluton that grew episodically since ca. 390 Ma during voluminous melt generation periods within the middle to lower crust, which also resulted in voluminous volcanism. The geochemical trends of the felsic volcanic rocks reflect the entrainment of xenoliths of peritectic garnet, cordierite and feldspar, and as such, the garnet-bearing felsic volcanic rocks represent an erupted mixture of a lower-temperature (*ca.* 770 °C) silicic melt and autocrysts, and peritectic phases and zircon crystals from previously crystallized and stored felsic melts.

Keywords Peritectic garnet · Prograde monazite · Peraluminous rhyolites · Petrogenesis · Iberian Pyrite Belt

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Introduction

Crustal-derived water-undersaturated and peraluminous felsic plutonic and volcanic rocks, broadly corresponding to S-type granites of Chappell and White (1992), represent a major component of the upper continental crust. They are typically generated by partial melting of a metasedimentary crust under variable temperature and pressure conditions, involving the breakdown of muscovite and biotite (Clemens et al. 2020; Johnson et al. 2021).

Experimental work and petrology studies on upper amphibolite- to granulite-facies metasedimentary migmatites and plutons in the lower and middle crust have provided deep insights into partial melting and melt extraction in the crust (e.g., Brown 2013; Clemens et al. 2020), whereby melt is generated via incongruent fluid-present or fluid-absent melting reactions. These commonly generate peritectic phases found in melt-depleted migmatites (e.g., Taylor et al. 2010; Rocha et al. 2017; Charette et al. 2021), which can be entrained and transported during segregation and extraction from the source (Taylor and Stevens 2010). Peritectic phases are not common in felsic plutons, as entrained peritectic phases may be rapidly assimilated into the melt (Clemens et al. 1997) or react with the host melt, generating other stable phases (Lavaure and Sawyer 2011; Rong et al. 2017). Nevertheless, numerous works have identified compositionally and texturally modified peritectic phases in felsic plutons (Erdmann et al. 2009; Villaros et al. 2009a; Taylor and Stevens 2010; Lavaure and Sawyer

2011; Dorais and Tubrett 2012; Jung et al. 2022; Dorais and Campbell 2023). The geochemical variation in major and trace elements of felsic plutonic and volcanic rocks has thus been increasingly interpreted as resulting from the entrainment and assimilation of peritectic phases (e.g., Stevens et al. 2007; Villaros et al. 2009b; Farina et al. 2012; Clemens et al. 2017b; Garcia-Arias and Stevens 2017; Zhu et al. 2020, 2021; Bailie et al. 2020).

The relationship between felsic plutons and volcanic rocks (the volcanic–plutonic relation), however, remains debated (e.g., Clemens et al. 2022). The most common view encountered in the literature is that felsic volcanic rocks represent melts (and their corresponding crystal cargo) extracted from crystallizing silicic crystal mushes (e.g., Bachmann et al. 2007; Wallrich et al. 2023). Other studies have shown that felsic melts can form and extrude from incrementally emplaced plutons without evidence of significant fractionation in crustal reservoirs, and preserve source-inherited geochemical signatures (e.g., Clemens et al. 2011a, 2017b; Fiannacca et al. 2017). The major challenge in linking the petrogenesis of felsic volcanic rocks to source-related processes is the scarcity of preserved peritectic phases (e.g., garnet) and granulite-facies xenoliths in the volcanic record, although they have been reported (Harangi et al. 2002; Cesare et al. 2009; Acosta-Vigil et al. 2010).

Garnet-bearing felsic volcanic rocks are found in the Cotovio area in the Iberian Pyrite Belt (IPB), the most important tectonostratigraphic unit of the South Portuguese Zone (SPZ) (Fig. 1). These garnet-bearing volcanic rocks are part of a suite of voluminous crustal-derived felsic magmas erupted in the IPB during the Paleozoic, but their

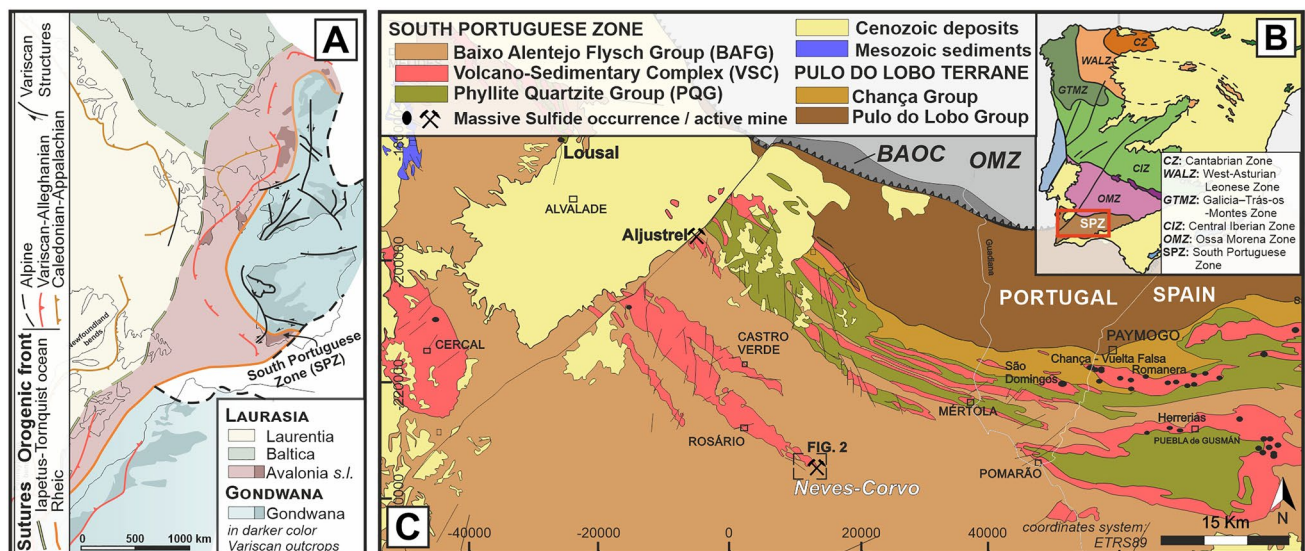


Fig. 1 A Simplified paleogeographic reconstruction showing the South Portuguese Zone before the fragmentation of Pangea (adapted from Pastor-Galán et al. 2020); B map of the Iberian Terrane (adapted

from Ribeiro et al. 2010); C geological map of the South Portuguese Zone, adapted from Luz et al (2021)

petrogenesis is yet to be fully understood. This work aims to identify the origin of garnet crystals and constrain the petrogenesis of these garnet-bearing felsic volcanic rocks, providing insights into the source to surface evolution of this crustal-derived magmatism.

Geological background

The closure of the Rheic Ocean during the Variscan orogeny culminated with the collision of Gondwana with Laurasia, whose suture can be found from Central America through Central Europe (Nance et al. 2012). In SW Iberia, the suture separates the Ossa Morena Zone (OMZ), the northern margin of Gondwana, and the SPZ on the Laurasian side (Fig. 1A; Ribeiro et al. 2007; Braid et al. 2011b).

The SPZ–OMZ collision developed under an oblique tectonic setting, generating transtensional pull-apart basins (Oliveira 1990; Quesada 1991; Tornos et al. 2002), which constitute one of the biggest volcanogenic massive sulfide (VMS) metallogenic districts in the world—the Iberian Pyrite Belt (IPB) (Tornos et al. 2002). These basins were infilled by a siliciclastic-dominated stratigraphic sequence

(up to 5–10 km thick), with ages ranging from the Middle Devonian up to the Carboniferous (Pereira et al. 2007; Mendes et al. 2020). These sediments were derived from eroded crustal basement, possibly the Meguma terrane (Braid et al. 2011, 2012; Luz et al. 2021). From the Viséan onward, the SPZ was deformed and metamorphosed under low-grade metamorphic conditions, up to lower-greenschist facies, generating a SW-verging thin-skinned fold–thrust belt (Munhá 1990; Abad et al. 2001).

Crustal-derived felsic volcanic rocks (rhyolites to dacites) often dominate the bimodal volcanism in the IPB (Mitjavila et al. 1997; Tornos 2006), although mafic volcanic and subvolcanic intrusive rocks can be voluminous in some sectors. Intermediate clinopyroxene-bearing volcanic rocks also crop out in the IPB and formed by crustal contamination and/or mixing of mantle-derived mafic magmas with crustal-derived felsic melts (Mitjavila et al. 1997; Codeço et al. 2018; Donaire et al. 2020a). U–Pb age dating of the felsic volcanic rocks in the IPB indicate that magmatism spanned from ca. 380 to 335 Ma (Rosa et al. 2009; Oliveira et al. 2019; Albardeiro et al. 2023).

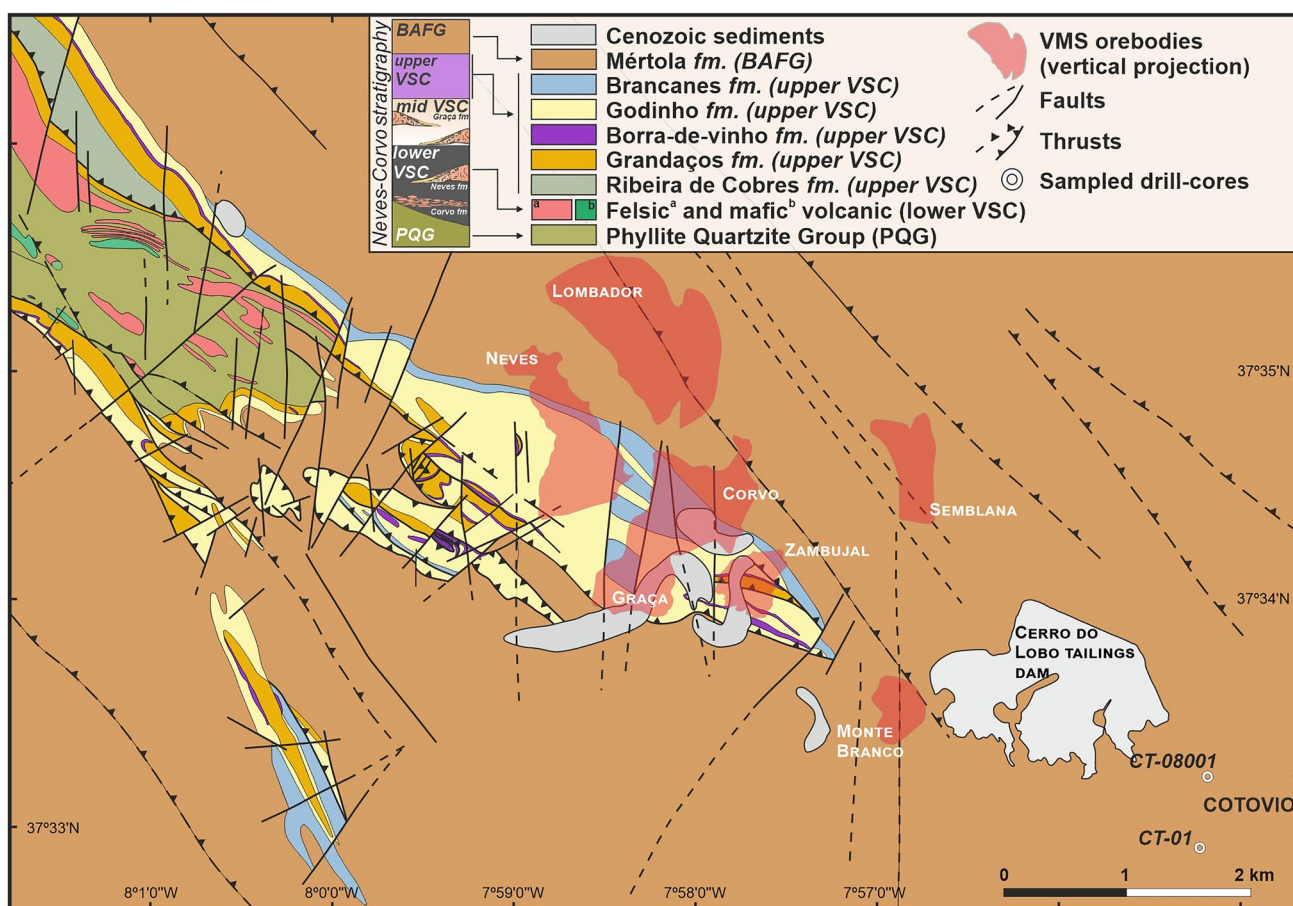


Fig. 2 Geological map of the Neves-Corvo area, adapted from Pereira et al. (2021) with a simplified lithostratigraphic column of the area

The Rosário–Neves–Corvo anticline (Figs. 1C, 2) represents a SW-verging major Variscan structure in the SPZ (Oliveira et al. 2004, 2013). The anticline gently plunges to the SE, and associated deformation comprises intense foliation near thrust faults with associated folding and moderate foliation development elsewhere in a thin-skinned deformation style. The stratigraphy in the Neves–Corvo area resembles the classical IPB stratigraphic sequence comprising from bottom to top: (i) the phyllite–quartzite group (PQG), of Givetian–Famennian age (as old as *ca.* 387–384 Ma), with abundant sandstones, graywackes, mudstones and marls (Oliveira et al. 2004; Mendes et al. 2020); (ii) the volcano-sedimentary complex (VSC), dominated by Late Famennian to Viséan (*ca.* 366–332 Ma) pelitic rocks (Oliveira et al. 2004; Pereira et al. 2021) with abundant felsic volcanic rocks of rhyolitic to dacitic composition, representing effusive felsic domes/cryptodomes and explosive volcanic pyroclastic deposits (Rosa et al. 2008; Oliveira et al. 2013); and (iii) the Baixo Alentejo Flysch Group (BAFG), deposited during the Viséan, and composed of siliciclastic flysch-type turbidites (Pereira et al. 2007). In the SE termination of the Rosário anticline, two mineral exploration drill holes intersected a thick garnet-bearing felsic volcanic unit in the hanging wall

stratigraphic sequence of the giant Sn-rich Cu–Zn Neves–Corvo VMS deposit (Carvalho et al. 2017) (Fig. 2).

Results

Volcanic facies and petrography

The garnet-bearing felsic volcanic rocks comprise thick and massive, variably muscovite- and chlorite-altered quartz–feldspar–phyric rhyolites (Fig. 3; Electronic Supplementary Material ESM 1). The volcanic sequence is dominated by the jigsaw-fit and clast-rotated breccia facies (Fig. 3). These consist of thick intervals of massive and poorly sorted clast-supported and clast- to matrix-supported breccias, respectively, with angular and blocky clasts set in a fine-grained matrix composed of quartz, feldspar and microphenocrysts fragments. Coherent intervals of microporphyritic and massive facies are less thick (up to a few meters) and are enclosed by the jigsaw-fit and clast-rotated breccia facies, with gradational contacts (Fig. 3). This volcanic facies association is typical of submarine felsic effusive lava flows and cryptodomes and is common throughout the IPB, where coherent volcanic

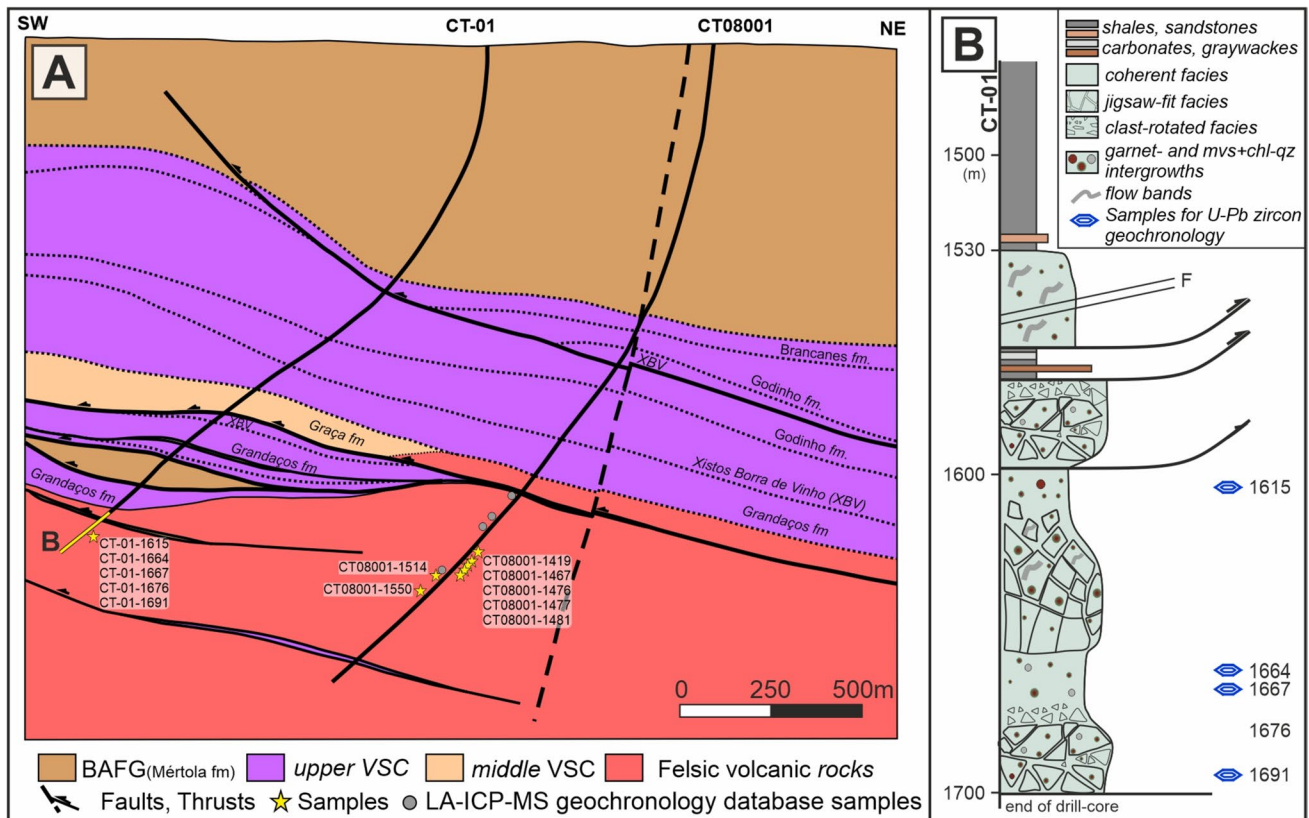


Fig. 3 **A** Geological cross section of the Cotovio area (adapted from Carvalho et al. 2017); **B** Detailed volcanic facies characterization of the CT-01 drill core

rocks are enveloped within hyaloclastites and thick in situ autobreccias, resulting from quenching and interaction with seawater, with minor sedimentary transport and reworking (e.g., McPhie et al. 1993; Rosa et al. 2008; Donaire et al. 2020b).

The matrix of the coherent volcanic rocks, devoid of fiamme, comprises micropoikilitic quartz feldspar (recrystallized volcanic glass) and muscovite, which also outlines the relict perlitic fractures (Fig. 4A, B). Phenocrysts account for up to 15%_{vol}. Quartz phenocrysts are common and are up to 0.5 mm in size (average 100 μm). They have rounded and embayed anhedral shapes (Fig. 4B–D), which display undulose extinction and are commonly mantled by fine-grained quartz. Variably sized (up to 4 mm) euhedral–subhedral porous and “dusty” albitized (?) feldspars are also abundant, and variably replaced by epidote (Fig. 4A–E) and muscovite. Epidote is also found as rounded crystal aggregates, replacing spherulites and the felsic matrix. Quartz and feldspars, although commonly found as individual microphenocrysts, are also found as rounded to subhedral crystal aggregates and as glomerocrysts. Biotite is rare, occurring as chloritized/ altered rounded microphenocrysts dispersed within the groundmass, exhibiting oxide-rich (opacitic) rims (Fig. 4C). Euhedral zircon crystals are common inclusions in feldspars (Fig. 4A), and are also found within the groundmass. Other accessory mineral phases include apatite and monazite, both disseminated in the matrix and as inclusions in feldspars. Primary oxide phases in the matrix and as inclusions in feldspars are often intensely replaced by TiO₂ (rutile) + titanite ± chlorite (ESM1 Sfigs 1.3N). Rare sulfides (pyrite ± chalcopyrite ± sphalerite) are found within the matrix and associated with apatite + chlorite + epidote + monazite + sulfides and carbonate veinlets and fractures. The matrix in the clast-rotated and jigsaw-fit breccias resembles the matrix in the coherent volcanic facies and is composed of abundant microcrystalline quartz and feldspar with fragments of quartz and feldspar microphenocrysts, but it is more fine-grained, and muscovite and chlorite are more abundant. Regardless of the volcanic facies, variably sized garnets are common (Figs. 4, 5; ESM1).

Garnet textures

Garnet in the felsic volcanic rocks is found dispersed in the matrix as poikiloblastic garnet–quartz intergrowths (Figs. 4, 5 Electronic Supplementary Material 1), quartz-free euhedral to subhedral garnet crystals (Fig. 5), or as crystal fragments.

Poikiloblastic garnet–quartz crystal intergrowths are sub-rounded to subhedral, sometimes with well-developed garnet crystal faces (Fig. 4F, G, H) and are up to 0.5 cm diameter. These garnets are characterized by abundant

elongated, filmy, rounded, euhedral, lobate to cusped quartz intergrowths and inclusions (Figs. 4F–I, 5A, B). Rounded and irregular polycrystalline inclusions are found in the quartz intergrowths, possibly representing crystallized melt inclusions. Parts of the quartz intergrowths in garnet display optical continuity. Another feature of the garnet–quartz intergrowths is the common presence of amalgamated garnet–quartz interlocking crystal aggregates (Fig. 4G). Quartz-free garnet grains are smaller (up to 0.1 cm diameter) and display subhedral to euhedral shapes (Fig. 5C–G).

Garnets contain small (< 40 μm) inclusions of euhedral–subhedral zircon and monazite (Figs. 4, 5), and occasionally inclusions of small aggregates (pseudomorphs?) of rutile, titanite and/or minor chlorite (ESM1.3, Sfigs. B, F, I, J, M), which sometimes also have apatite and quartz inclusions. Inclusions occur in the core, mantle and near the rims of the garnet–quartz intergrowths and quartz-free euhedral garnets (Figs. 4, 5; ESM1). Furthermore, garnet crystals are variably replaced by chlorite or chlorite and muscovite (Figs. 4, 5C, E, F). In zones where garnets are intensely replaced, abundant rounded and irregularly shaped monazite is sometimes found, with occasional minor allanite, rutile, and apatite.

Other rounded muscovite–chlorite–quartz intergrowths are also rarely found (Fig. 5H, I). They are texturally similar to the garnet–quartz intergrowths in size (up to 0.3 cm), comprising rounded, lobate, and cusped quartz, partially in optical continuity (Fig. 5H, I), intergrown with a parent phase which is completely altered to fine-grained aggregates of a muscovite and chlorite rich pseudomorph (pinite). These occur as complex replacements along microfractures that grew inward into the pseudomorphed phase (cordierite?) (Fig. 5H, I; ESM1). Accessory phases such as zircon, monazite, apatite and minor allanite are also found within the pseudomorphous aggregates (Fig. 5H).

Garnet major and trace element composition

The complete garnet EPMA and LA-ICP-MS dataset can be found in ESM2 Table 1, additional LA-ICP-MS trace-element profiles in ESM3, and analytical methods are provided in ESM4. All garnets are FeO-rich and MgO-, MnO-, and CaO-poor, almandine-pyrope solid solution garnets, with minor spessartine and grossular components (between Alm_{0.90–0.80}Pyro_{0.10–0.04}Sps_{0.05–0.01}Grs_{0.03–0.01}) (EPMA data in ESM2 Table 1). Garnets intergrown with quartz show major element zoning, with preserved normal prograde growth zoning characterized by rimward increases in X_{Alm} (0.85 up to 0.90) and X_{Pyro} (0.05 up to 0.06) and decreases in X_{Sps} (0.03 down to 0.01) and X_{Grs} (0.02–0.01) (Fig. 6A, B). Quartz-free, euhedral garnets are characterized by similar compositions, but differ from the garnet intergrown with quartz by having lower X_{Sps}

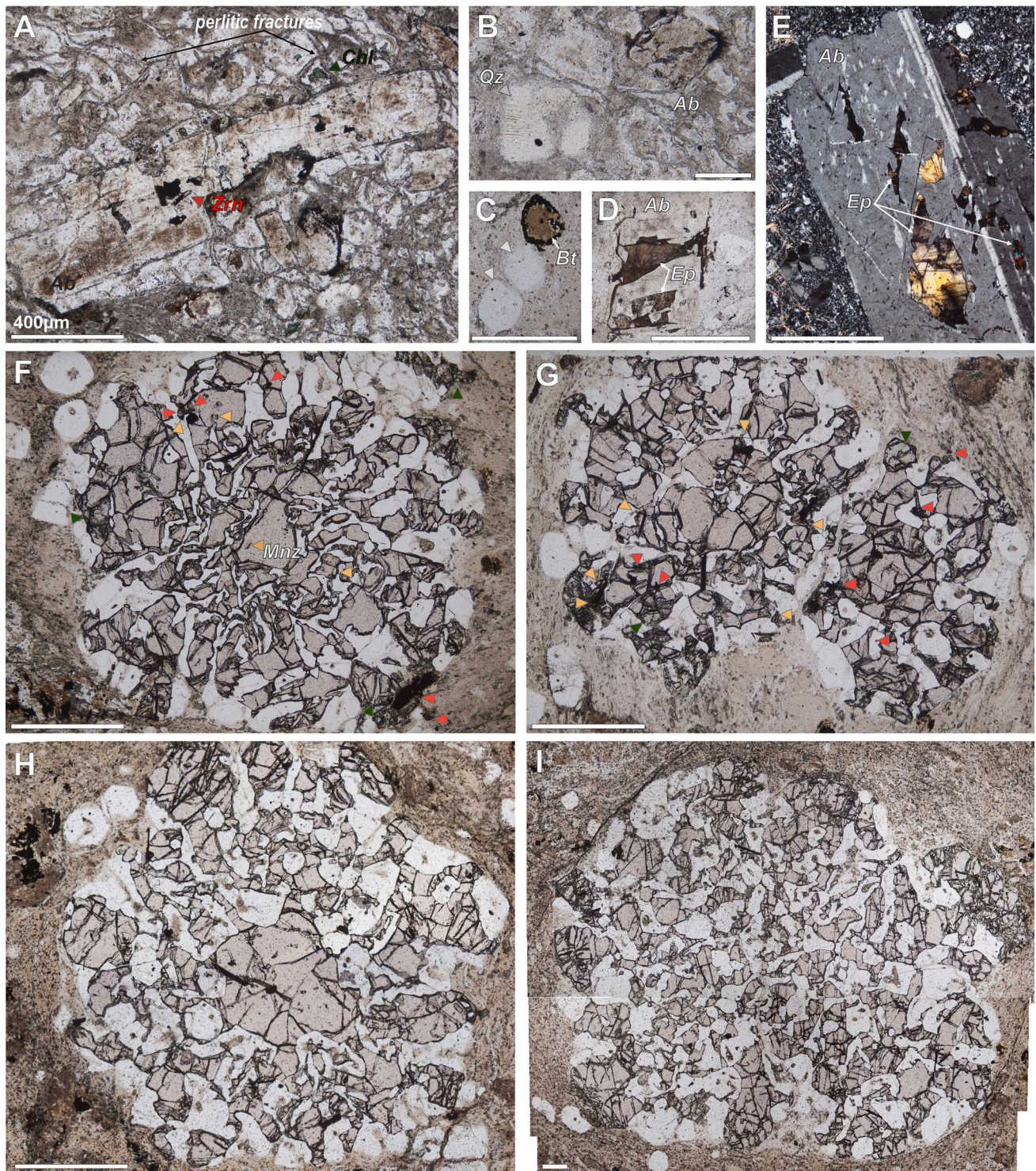


Fig. 4 Photomicrographs of the garnet-bearing silicic volcanic rocks. Red arrows represent zircon inclusions, orange arrows are monazite inclusions in garnet and quartz, and dark-green arrows are chlorite. Scale bars are 400 μm . **A** Zircon inclusions in a feldspar phenocryst (albite), set in a quartz- and K-feldspar-rich matrix with abundant perlitic fractures; **B** Embayed quartz and euhedral feldspar

microphenocrysts. **C** Rounded biotite with opaque rims and quartz microphenocrysts. **D** and **E** Epidote replacing feldspars, **F** to **I** rounded to subhedral garnet intergrown with quartz, displaying filmy, elongated and lobate quartz inclusions. Note in **G** the presence of two amalgamated garnet–quartz intergrowths

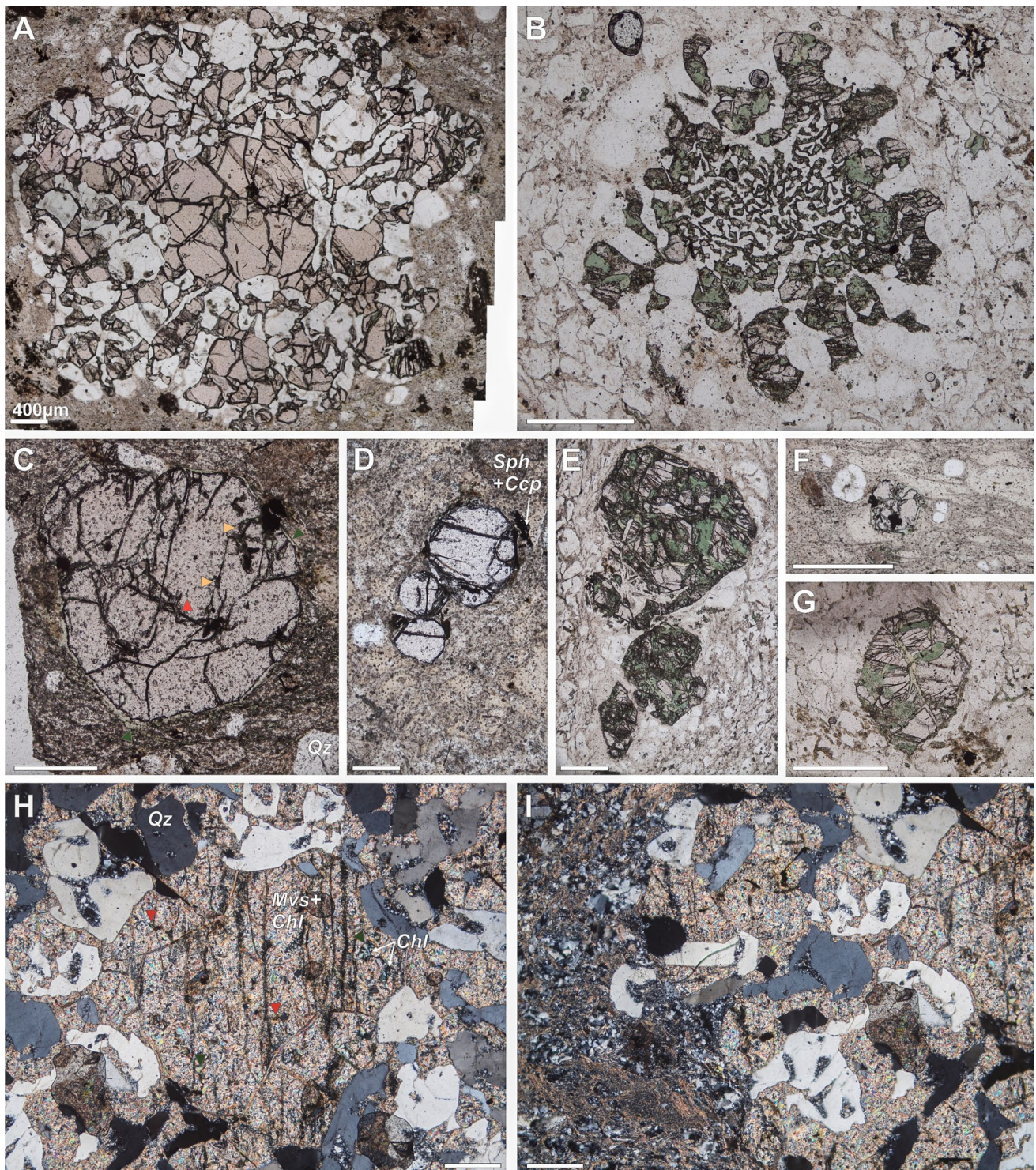


Fig. 5 Photomicrographs of the garnet-bearing silicic volcanic rocks. Red arrows are zircon inclusions, orange arrows are monazite inclusions in garnet and quartz and green arrows are chlorite. Scale bars are 400 μm . **A** Garnet intergrown with quartz displaying a quartz-free core and filmy and lobate quartz inclusions and intergrowths in the mantle and rim (trace element map in Fig. 7). **B**

Chlorite-altered garnet intergrown with lobate quartz intergrowths, **C** to **G** smaller, euhedral quartz-free garnets, variably replaced by chlorite. **H** and **I** Muscovite–chlorite pseudomorphs intergrown with quartz. Note that quartz is not entirely in optical continuity, a similar feature to the garnet–quartz intergrowths and inclusions

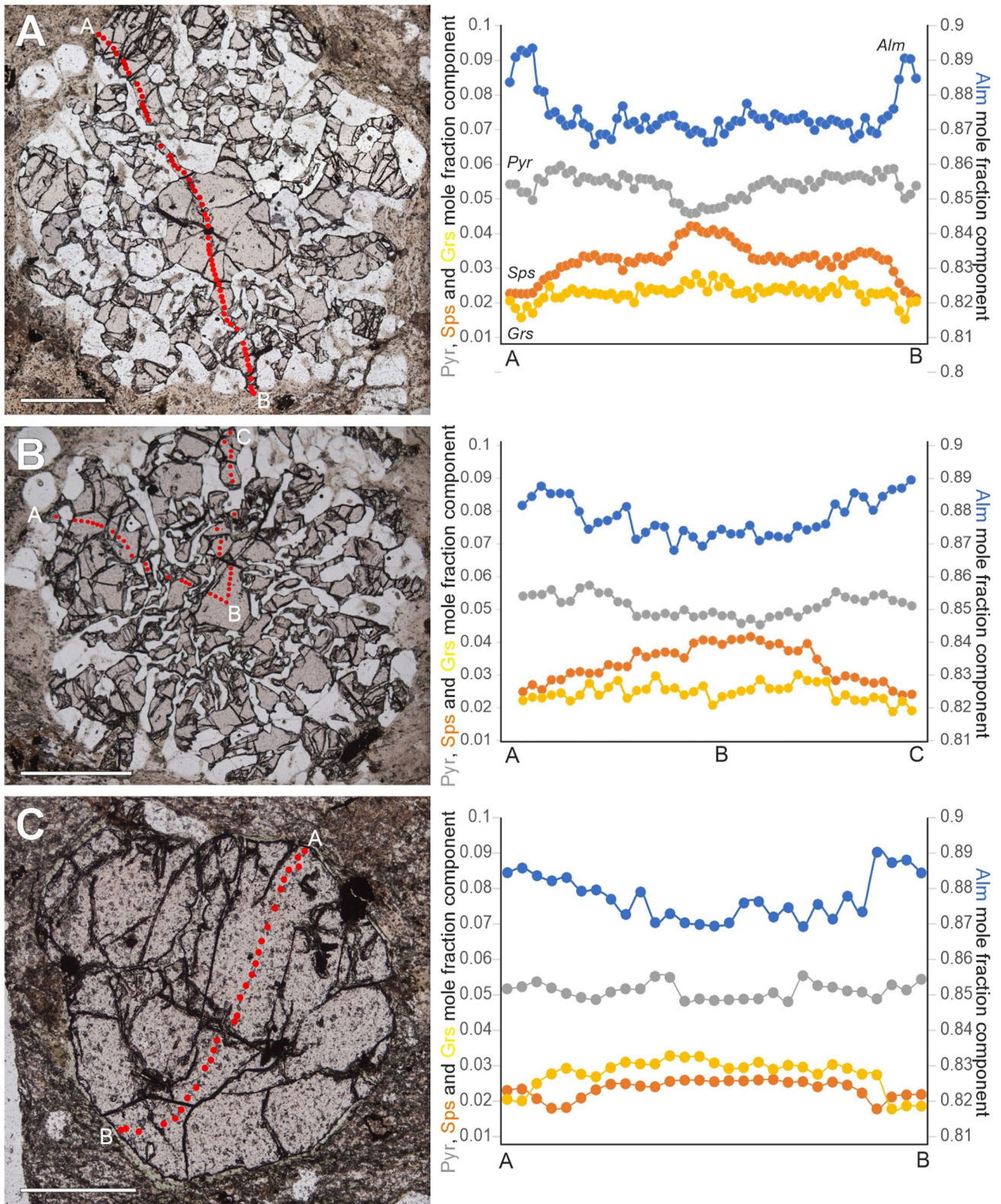


Fig. 6 A–C Representative rim-to-rim EPMA major element profiles of two garnets intergrown with quartz and one smaller euhedral garnet. Scale bars are 400 μm

and comparatively higher X_{Grs} (Fig. 6C). Euhedral garnets often exhibit no major element zoning, although larger sized crystals display some major element chemistry zoning, with Ca- and Mn-enriched ($X_{\text{Grs}} > X_{\text{Sp}}^{\text{e}}$) cores and slight rimward increases in Mg and Fe (Fig. 6C). Both garnet types occasionally show the presence of thin retrograde rims in contact with the groundmass, with small increases in Mn and Mg and similar decreases in Fe (Fig. 6C).

The mapped garnet intergrown with quartz (Fig. 7) shows a decrease in Mn toward the rim and a small increase in the mantle area. Furthermore, there is prominent concentric trace element zoning, characterized by the presence of an inner core enriched in V, Sc, HREE, Zr and Ti, and an outer core marked by a sharp decrease in trace element concentrations (Fig. 7). Both core domains have well-developed oscillatory zoning, as best displayed in the V and HREE maps (Fig. 7). The outer core is overgrown by a trace element-enriched mantle (HREE, Sc, V, Ti and Zr), with marked oscillatory zoning, and rimward trace element decreases (Fig. 7). The trace element-rich mantle is broader in size in the HREE, Sc and V maps, whereas the Zr and Ti maps display a thinner mantle (Fig. 7), although these differences can also be related to the thickness of the oscillatory zoning. Furthermore, the outer core–mantle transition seems to coincide with the appearance of filmy, lobate quartz intergrowths, but there is no evidence of dissolution between the core–mantle and mantle–rim domains. Also noticeable in some EPMA profiles in Fig. 6, the mantle area shows a slight increase in Mn (Fig. 7) and Ca (ESM3).

Garnets intergrown with quartz display a wide range in Y (167–3250 $\mu\text{g/g}$), P (119–733 $\mu\text{g/g}$), $\sum\text{REE}$ (133–1773 $\mu\text{g/g}$) and $\sum\text{HREE}$ (121–1758 $\mu\text{g/g}$), whereas smaller euhedral garnets have markedly lower trace element concentrations (Y 171–1486 $\mu\text{g/g}$, P 186–383 $\mu\text{g/g}$, $\sum\text{REE}$ 133–1136 $\mu\text{g/g}$ and $\sum\text{HREE}$ 121–856 $\mu\text{g/g}$). Ti concentrations varies from 152 to 1870 $\mu\text{g/g}$ in garnet intergrown with quartz, and 408 to 1061 $\mu\text{g/g}$ for euhedral garnets, while Zr and Hf concentrations are overall similar in both euhedral garnets and garnets intergrown with quartz (20–260 $\mu\text{g/g}$ and 30–250 $\mu\text{g/g}$, respectively). Chondrite-normalized REE patterns of all garnet analyses show strong LREE depletion and M-HREE enrichment, variably pronounced Eu anomalies (Eu/Eu* between 0.02 and 0.07) and highly variable $\text{Lu}_\text{N}/\text{Gd}_\text{N}$ ratios (Fig. 8). Garnet intergrown with quartz shows wider ranges in the $\text{Lu}_\text{N}/\text{Gd}_\text{N}$ ratios (from 0.03–8.09) than the euhedral quartz-free garnet (0.06–0.73) (Fig. 8). LA-ICP-MS trace element profiles from other garnet–quartz intergrowths show Y + HREE rich cores and a variably trace element enriched mantle, with sharp decreases in trace element concentrations between both domains (ESM3). This zoning is particularly noticeable in Y, Lu and Yb in all the garnet transects, whereas MREE,

Ti, Zr, P and Hf display wide variations in the core and mantle zones (ESM3). A trace element profile (rim-to-rim) of a euhedral garnet shows the presence of an inner core with high Zr, Ti, Hf, P and MREE, and low HREE + Y (Fig. 8). The sharp decrease in Zr, Ti and MREE in the outer core area coincides with increases in Y, Yb and Lu, whereas Y and HREE decrease toward the rim with the trace element profile showing a second mantle enrichment in Zr, Ti, MREE and P (Fig. 8).

Whole-rock geochemistry

Major and trace element compositions of the garnet-bearing volcanic rocks are presented in ESM2 Table 3 and methods are in ESM4 (Fig. 9). All samples show moderate to high SiO_2 contents (67.2–80.1 wt%), with $A/\text{CNK} > 1$, mostly above 1.1. Most major elements and Zr define scattered and broad negative correlations with increasing SiO_2 (Fig. 9; ESM2 Table 3). $\text{FeO}_\text{T} + \text{MgO}$ (maficity), Al_2O_3 and TiO_2 as well as Eu, U and Zr show quasi-linear negative correlations with SiO_2 (Fig. 9). A/CNK , TiO_2 , Al_2O_3 , Mg# define positive trends with increasing $\text{FeO}_\text{T} + \text{MgO}$, as well as Zr, Hf, Yb, Sm and Nb, although with some scatter, whereas CaO and SiO_2 display negative correlations with $\text{FeO}_\text{T} + \text{MgO}$ (Fig. 9).

The garnet-bearing felsic volcanic rocks are also characterized by variable, but overall high total REE concentrations (41–431 $\mu\text{g/g}$) and markedly negatively sloped C1-normalized REE patterns ($\text{La}_\text{N}/\text{Yb}_\text{N}$ from 3.6–33.1). REE patterns have negative LREE and HREE slopes ($\text{La}_\text{N}/\text{Sm}_\text{N}$ and $\text{Gd}_\text{N}/\text{Lu}_\text{N}$ between 1.13–3.4 and 1.2–5.9), negative Eu anomalies ($\text{Eu}/\text{Eu}^* = \text{Eu}_\text{N}/(\text{Sm}_\text{N} \times \text{Gd}_\text{N})^{0.5}$) of 0.28 to 0.40 (0.33 average) and mildly positive Ce anomalies (0.92–1.14, avg. 1) (Fig. 10A). Primitive mantle (PM)-normalized trace element diagrams show pronounced enrichments in LILE (e.g., Rb, Ba), Th and U over HSFE (e.g., Nb), with marked negative anomalies in Pb, Sr and Ti (Fig. 10B). Moreover, the garnet-bearing felsic volcanic rocks show low Nb (8–24 $\mu\text{g/g}$), Ta (0.8–1.6 $\mu\text{g/g}$) and sub-chondritic Nb/Ta (9–18), low Cr (< 0.5–8.3 $\mu\text{g/g}$), V (< 5 $\mu\text{g/g}$), and Ni (2–9 $\mu\text{g/g}$), and Zr concentrations between 80 and 315 $\mu\text{g/g}$.

Zircon morphology, geochronology and geochemistry

All dated samples from both drill cores represent different facies of a specific volcanic deposit, as indicated by geological, stratigraphic, and physical volcanology data. Thus, U–Pb zircon data are integrated with geological and stratigraphic data, providing a robust framework with which to interpret the results. Zircon populations were calculated using the Sambridge and Compston (1994) method. All

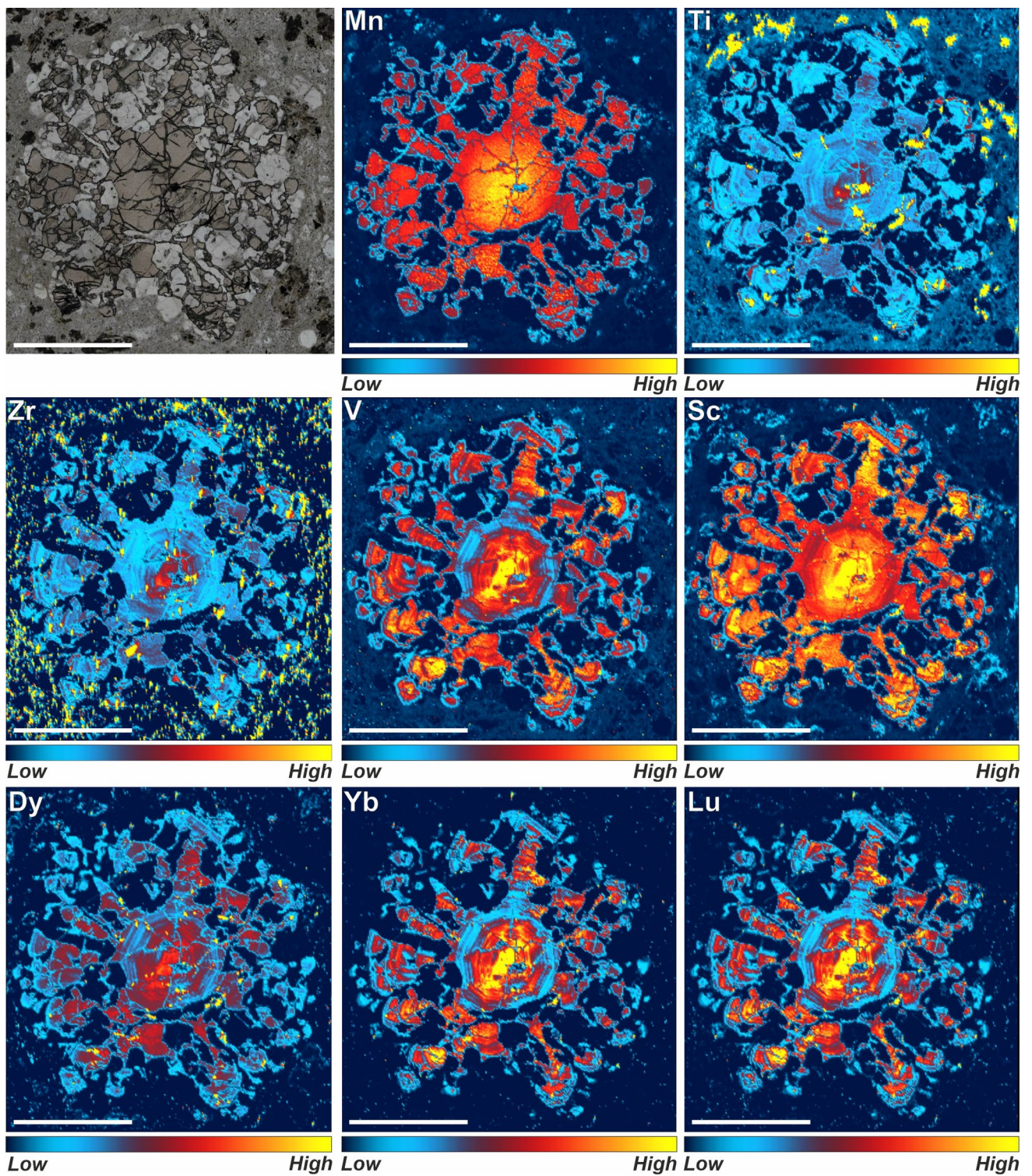


Fig. 7 LA-ICP-MS trace element maps of a garnet intergrown with quartz. Scale bars are 400 μm. Additional maps of Ca, Ce, Gd and Dy and trace element spot analyses profiles of garnet–quartz intergrowths are provided in ESM3

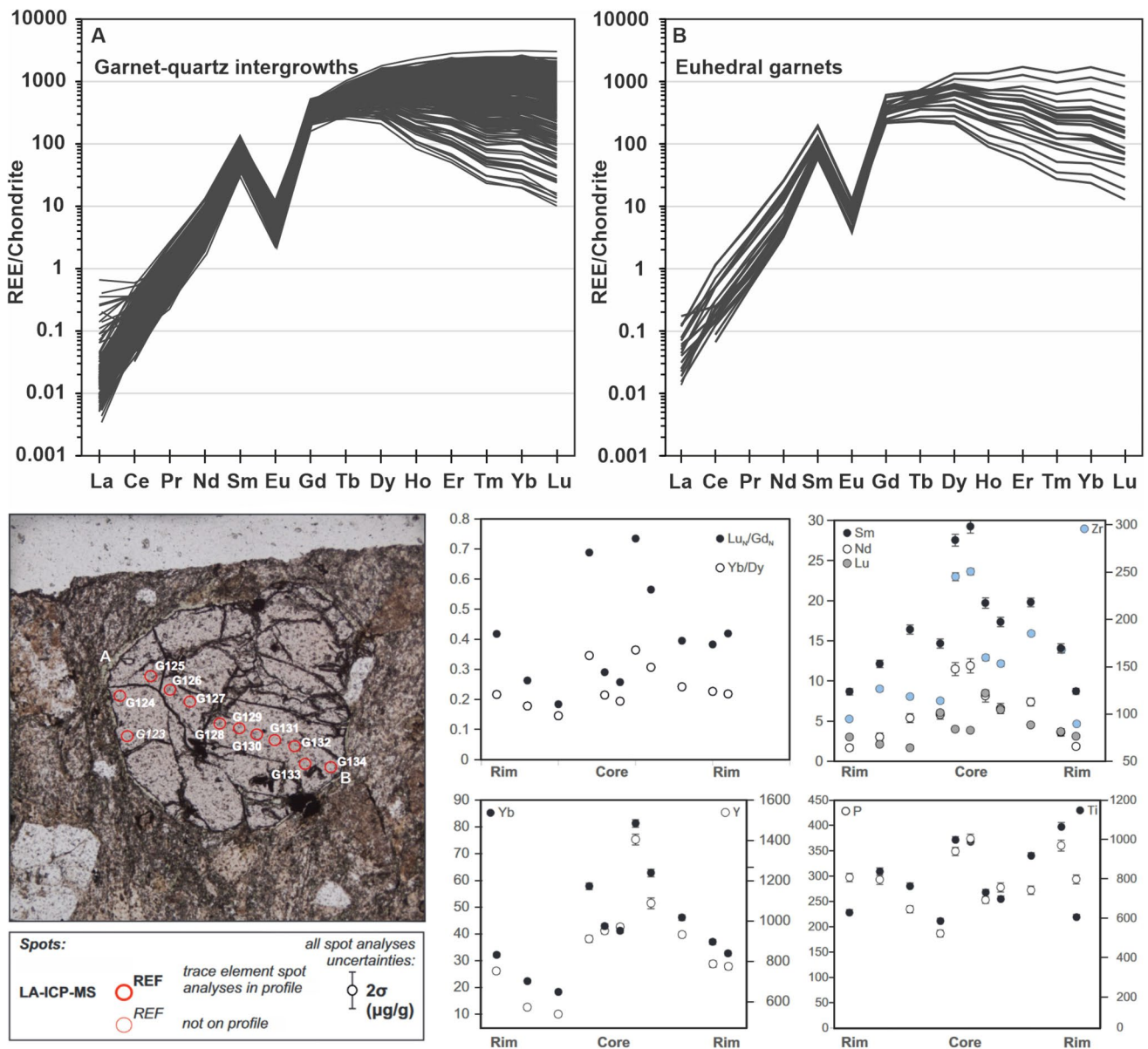


Fig. 8 **A** and **B** Chondrite-normalized garnet REE data [values from McDonough and Sun 1995]; **C** Trace element (Zr, REE, Y, P and Ti) variations along a rim-to-rim profile **A–B** for a euhedral garnet (major element in Fig. 6)

reported uncertainties for individual analyses and calculated zircon populations are 2σ. Data are provided in ESM2 Table 5 and 6 and detailed methods are provided in ESM4.

SHRIMP U–Pb zircon geochronology

Zircon from the four CT-01 drill core samples comprise small (<200 μm in length) euhedral to subhedral grains or fragments, and are transparent, prismatic, and elongated (from 2:1 to 4:1 length/width ratios). All the grains show homogeneous textures, fractures and the presence of inclusions in BSE images. Cathodoluminescence (CL)

images show some variation of CL response and the common presence of typical magmatic textural features (as in Corfu et al. 2003) such as oscillatory zoning and sector zoning, and in some grains a homogeneous CL response (ESM1). Most grains are characterized by similar morphologies, with well-developed {211} pyramids, dominating over {101}, and dominant {110} prisms over {100}. There is no relationship between specific textural features or CL response to any of the individual zircon populations. One important feature is the lack of overgrowths and texturally complex zircons.

Zircon U–Pb dates from all four samples (CT-01 drill-core) display a wide range of concordant dates (Fig. 11A,

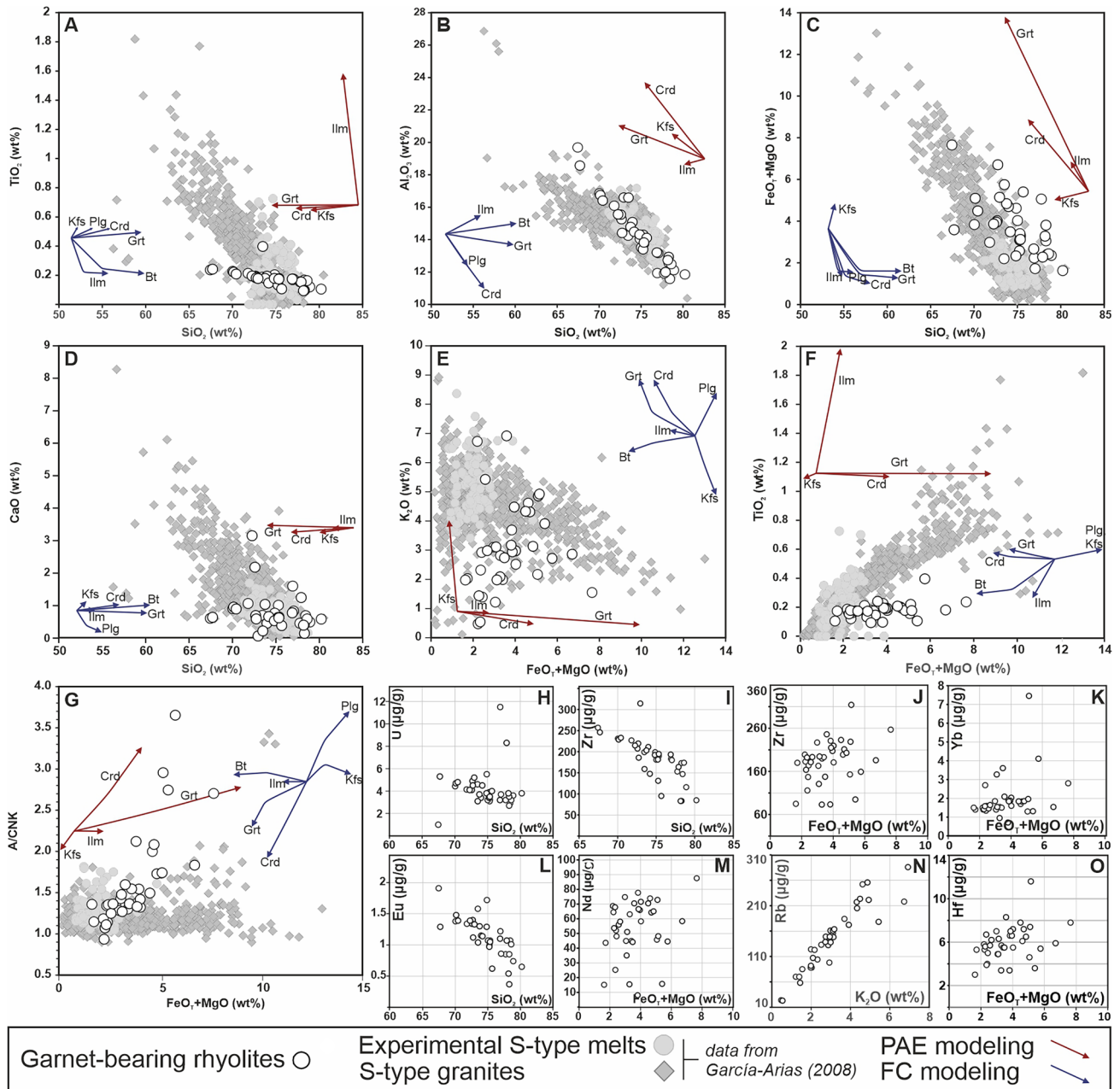


Fig. 9 A to G Major element Harker plots for the Cotovio garnet-bearing rhyolites (recalculated to a volatile-free basis). A TiO_2 ; B Al_2O_3 ; C $\text{FeO}_T + \text{MgO}$; D CaO. E K_2O vs $\text{FeO}_T + \text{MgO}$; F TiO_2 vs $\text{FeO}_T + \text{MgO}$; G A/CNK vs $\text{FeO}_T + \text{MgO}$; H U vs SiO_2 ; I Zr vs SiO_2 ; J Zr vs $\text{FeO}_T + \text{MgO}$; K Yb vs $\text{FeO}_T + \text{MgO}$; L Eu vs SiO_2 ; M Nd vs $\text{FeO}_T + \text{MgO}$; N Rb vs K_2O ; O Hf vs $\text{FeO}_T + \text{MgO}$.

Additional information on peritectic assemblage entrapment (PAE) and fractional crystallization (FC) modeling are provided in ESM2 Table 4. Plots A–G show also the vectors calculated for peritectic phase entrapment (in red), of garnet, cordierite, K-feldspar (20%) and ilmenite (2%), and fractional crystallization (in blue) of plagioclase, K-feldspar, biotite, garnet and cordierite (10%), and ilmenite (2%)

C), suggesting the presence of multiple zircon populations. The older $^{206}\text{Pb}/^{238}\text{U}$ dates ($n=4$), from all samples, scatter between $ca. 470 \pm 12$ Ma up to $ca. 439$ Ma, with variable Th/U ratios (0.12–1.98; Fig. 11E), whereas the remaining $^{206}\text{Pb}/^{238}\text{U}$ dates ($n=47$, from all samples) range from $ca.$

400 Ma up to $ca. 351$ Ma. The analyses define four main zircon populations at 453 ± 6 Ma, 407 ± 5 Ma, 384 ± 6 Ma, and 365 ± 1.6 Ma (Fig. 11A), with Th/U ratios between 2.01 and 0.11 (Fig. 11E).

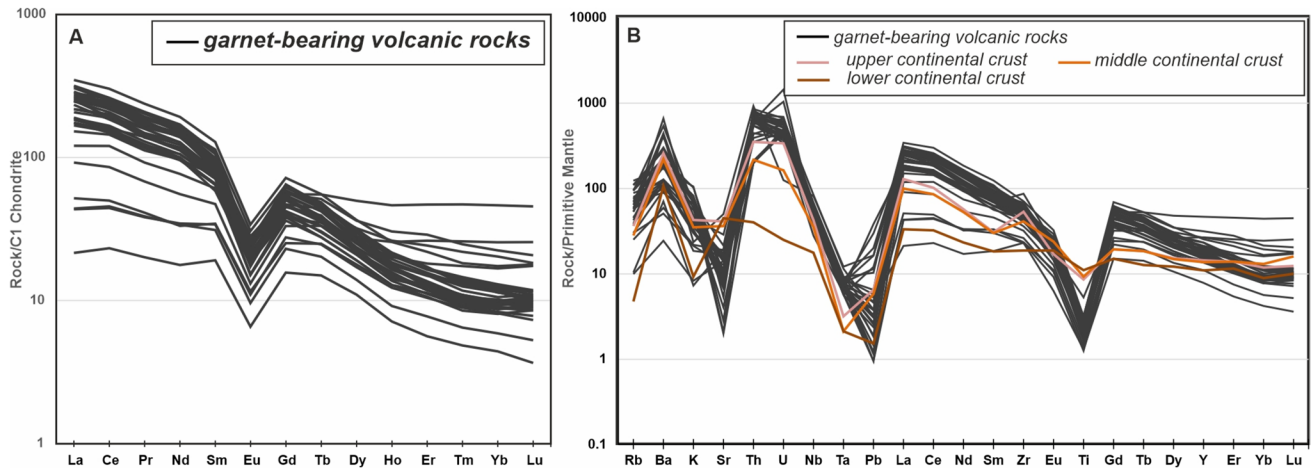


Fig. 10 **A** C1 Chondrite-normalized REE (values from McDonough and Sun 1995); **B** PM-normalized (values from Palme and O'Neill 2014) trace element plot of the garnet-bearing rhyolites whole-rock samples

LA-ICP-MS U–Pb zircon geochronology

Zircon grains from the four LA-ICP-MS dated samples of the CT08001 drill core are small (<200 μm in length). The euhedral zircon grains and zircon fragments with length–width ratio of 2:1 to 4:1. Common CL textures include oscillatory zoning or a homogeneous CL response. Zircons also show well-developed {211} pyramids and {110} prisms, dominating over {101} pyramids and {100} prisms.

The zircon LA-ICP-MS U–Pb data of all four samples also show a wide variation in concordant $^{206}\text{Pb}/^{238}\text{U}$ dates ($n=104$ analyses within 10% concordance) spanning in a multi-peak age spectrum (Fig. 11B). The youngest dates ($n=69$, from all samples) range from 338 ± 4 up to 378 ± 6 Ma, and comprise most of the data. They define four major peaks at 345 ± 1 Ma, 359 ± 1 Ma, 366 ± 1 Ma, and 374 ± 1 Ma. These overlap with the main volcanic episodes in the stratigraphy in Neves-Corvo (Fig. 11B), whereas the remaining older $^{206}\text{Pb}/^{238}\text{U}$ dates ($n=35$, from all samples) range from 387 ± 6 and 453 ± 6 Ma, defining several minor peaks between 390 and 400 Ma (Fig. 11B).

Zircon trace element geochemistry

The LA-ICP-MS zircon trace element data is presented in ESM2 Table 7 and the analytical methods in ESM4. The zircon crystals from all individualized populations display coherent REE patterns (Fig. 12A), characterized by strong MREE and HREE enrichment, variable MREE–HREE segments ($\text{Yb}_\text{N}/\text{Dy}_\text{N}$ between 0.68 and 6.65), low $\text{Gd}_\text{N}/\text{Yb}_\text{N}$ (0.03–0.67) and $\sum\text{LREE}$ (0.44–47.78 $\mu\text{g/g}$), and variable negative Eu and positive Ce anomalies (0.02–0.7

and 0.83–4.89, respectively; Fig. 12). Noticeably, a few zircon analyses with ages from *ca.* 350–370 Ma (defining a *ca.* 365 Ma main population) display lower $\text{Gd}_\text{N}/\text{Yb}_\text{N}$ values and flat REE patterns (Fig. 12A). Other trace element compositional ranges of the zircon populations are also similar, with variable Hf (9560–15100 $\mu\text{g/g}$), Y (1079–11400 $\mu\text{g/g}$), Ti (3.42–34.2 $\mu\text{g/g}$, two outliers of 59 and 72 $\mu\text{g/g}$) concentrations (Fig. 12B–E). The majority of the zircon grains show LREE-I values ($\text{Dy}/\text{Nd} + \text{Dy}/\text{Sm}$) above 10, indicating no significant post-crystallization alteration (Bell et al. 2019). Ti is positively correlated with Th/U (Fig. 12D), and negatively correlated with Hf (Fig. 12F), the latter suggesting decreasing temperatures with magmatic differentiation. Hf also displays a positive correlation with Y (Fig. 12E), and the data appear to define two main trends, one with higher Y (above 9000 $\mu\text{g/g}$) and one with lower (up to 6000 $\mu\text{g/g}$) Y contents. Additionally, Y shows a broad and scattered negative correlation with Ti, also displaying two main compositional trends (Fig. 12H). Furthermore, Y is positively correlated with Gd/Yb ratios. Eu anomalies show a positive correlation with Hf (Fig. 12G). The overall low positive Ce/Ce* values suggest crystallization under reducing conditions, due to the higher compatibility of tetravalent Ce in relation to Ce^{3+} . Oxygen fugacity estimates, calculated using the zircon oxybarometer of Loucks et al. (2020) yield similar $f\text{O}_2$ estimates for all zircon analyses, mostly under reducing melt conditions and ΔFMQ values between 1.1 and –5.6. The average $f\text{O}_2$ of the *ca.* 365 Ma zircon population is -16.7 ± 2.1 (ESM3 Table 7). (Fig. 12, I, J). These results, although rather low, are within the range of $f\text{O}_2$ estimates for other reduced granites and silicic volcanic rocks (e.g., Pichavant et al. 1988, 2024; Bucholz et al. 2018; Zhu et al.

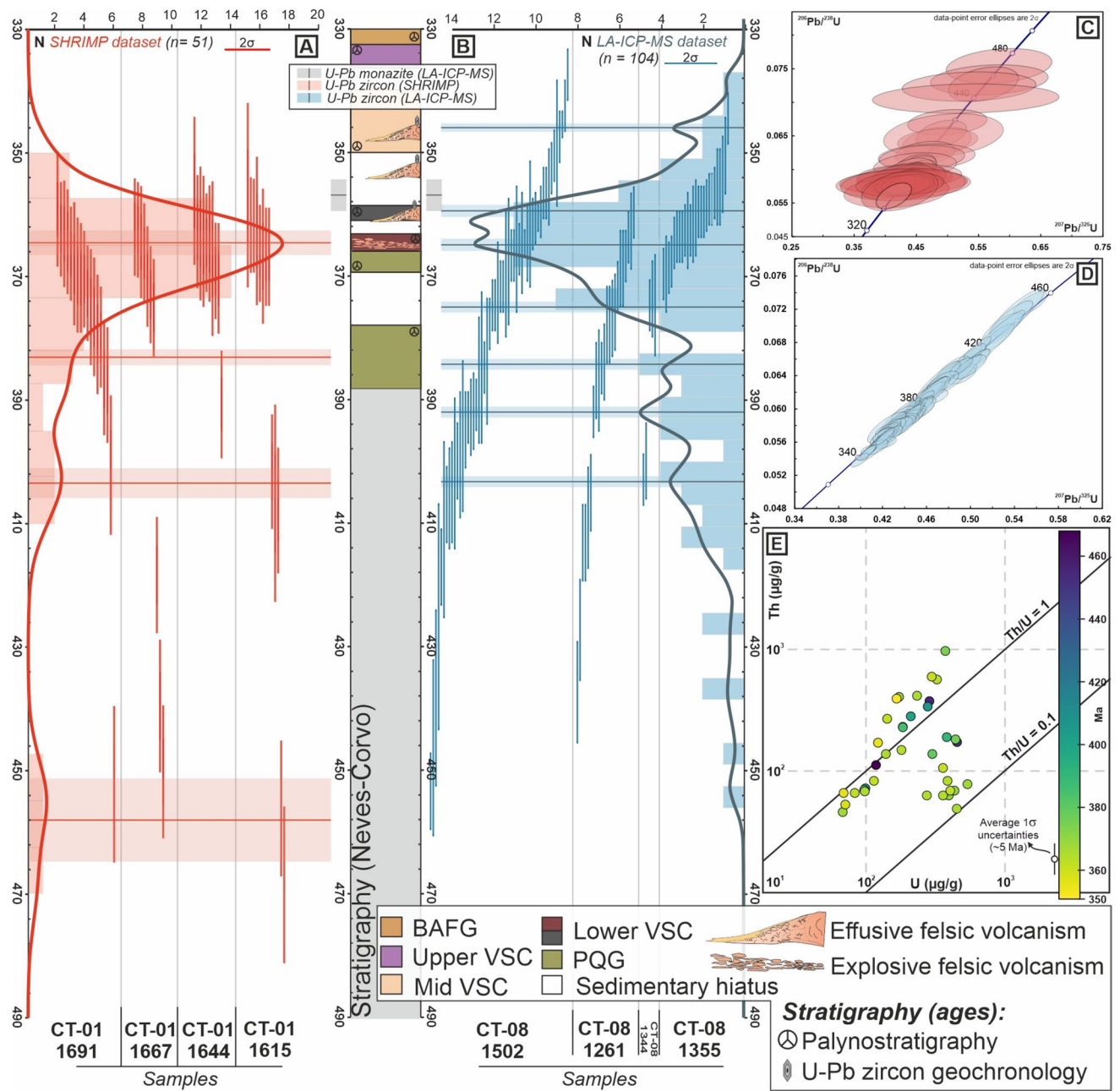


Fig. 11 **A** and **B** Probability density functions for the SHRIMP and LA-ICP-MS geochronology datasets with ranked U–Pb SHRIMP $^{206}\text{U}/^{238}\text{Pb}$ dates of the garnet-bearing felsic volcanic rocks; box heights are 2σ . Uncertainties in the individualized zircon populations

ages are calculated at the 95% (2σ) confidence level; **C** and **D** Concordia diagrams of the SHRIMP and LA-ICP-MS datasets; **E** Th vs U plot of the SHRIMP zircon analyses

2021; Li et al. 2021; Bell and Kirkpatrick 2021; Nazari-Dehkordi and Robb 2022; Sun et al. 2023; Huang et al. 2023; Vogt et al. 2023; Manor et al. 2023). Furthermore, there is a general trend of increasing ΔFMQ with Eu/Eu^* (Fig. 12F), suggesting the role of additional processes in controlling the Eu anomalies besides plagioclase fractionation (Holder et al. 2020; Bell and Kirkpatrick 2021).

Monazite geochemistry and in situ geochronology

Monazite geochemistry

Monazite EPMA data is presented in ESM2 and analytical methods in ESM4. There are four different monazite textural types with distinct compositions: (i) euhedral to

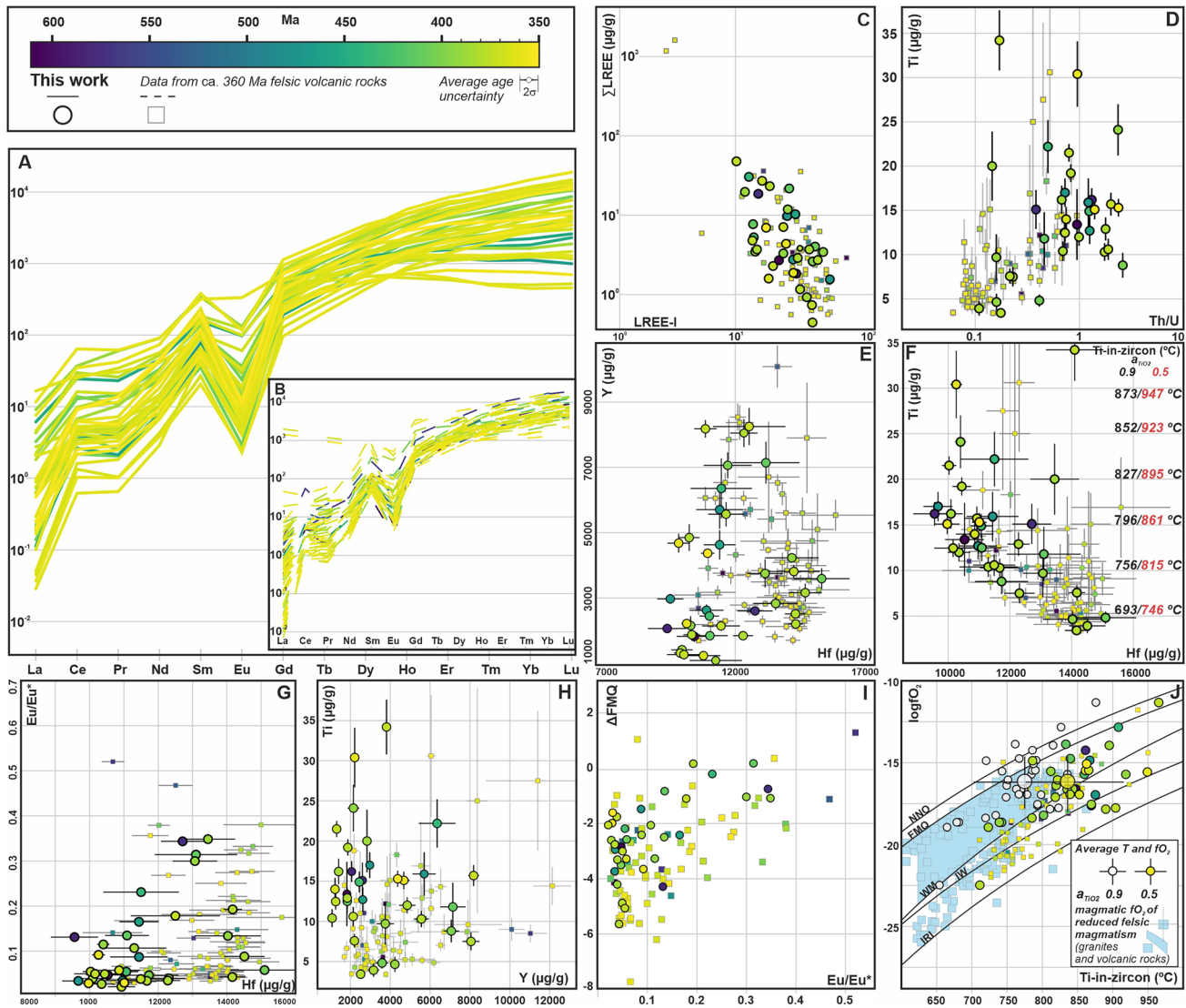


Fig. 12 Zircon trace element geochemistry **A** and **B** Chondrite C1-normalized (from McDonough and Sun 1995) zircon REE patterns; **C** LREE-I vs Σ LREE; **D** Ti vs Th/U; **E** Y vs Hf; **F** Ti vs Hf; **G** Eu/Eu* vs Hf; **H** Ti vs Y; **I** Δ FMQ vs Eu/Eu*; **J** $\log f_{O_2}$ vs $T_{\text{Ti-in-zircon}}$ ($^{\circ}\text{C}$). The plotted Ti-in-zircon and f_{O_2} average data is from

the ca. 365 Ma zircon population (see text for details). Data in blue (**J**) compiled from Li et al. (2021), Yang et al. (2000), Bucholz et al. (2018), Manor et al. (2023), Vogt et al. (2023) and Pichavant et al. (2024)

subhedral inclusions in garnet (Fig. 13A–H); (ii) inclusions in muscovite–chlorite pseudomorphs (Fig. 13I); (iii) inclusions in feldspars (Fig. 13J–K) and (iv) small (< 10 μm) anhedral grains near intense garnet replacements by chlorite and included in feldspars (Fig. 13L–M). All monazite analyses show X_{Mnz} -rich compositions (0.88–0.99; ESM2 Table 8), and their geochemical composition is controlled by the huttonite and minor cheralite substitutions (Fig. 13). Euhedral monazite inclusions in garnet and chlorite–muscovite pseudomorphs show variable Σ LREE and Y concentrations (YO_2 from 5.18–1.09 wt%, avg. 2.78 wt%), high Th ($\text{ThO}_2 > 2.88$ wt%) and low Si (apfu) (Fig. 13N–Q). These euhedral monazite inclusions are unzoned in Th and

Y, although some grains display some Y and Si variations (Fig. 13N–Q). Euhedral to subhedral inclusions in feldspar display textural and chemical features similar to the euhedral monazite inclusions in garnet and chlorite–muscovite pseudomorphs (Fig. 13N–Q). Anhedral monazite grains (rimming garnet and as feldspar inclusions) show the highest LREE contents, lower Th, and Y concentrations, and variable Si (apfu), up to 0.28 (Fig. 13N–Q).

Monazite U–Pb geochronology

Seventeen LA-ICP-MS U–Pb spots were obtained from four subhedral–euhedral Y- and Th-rich monazite inclusions in

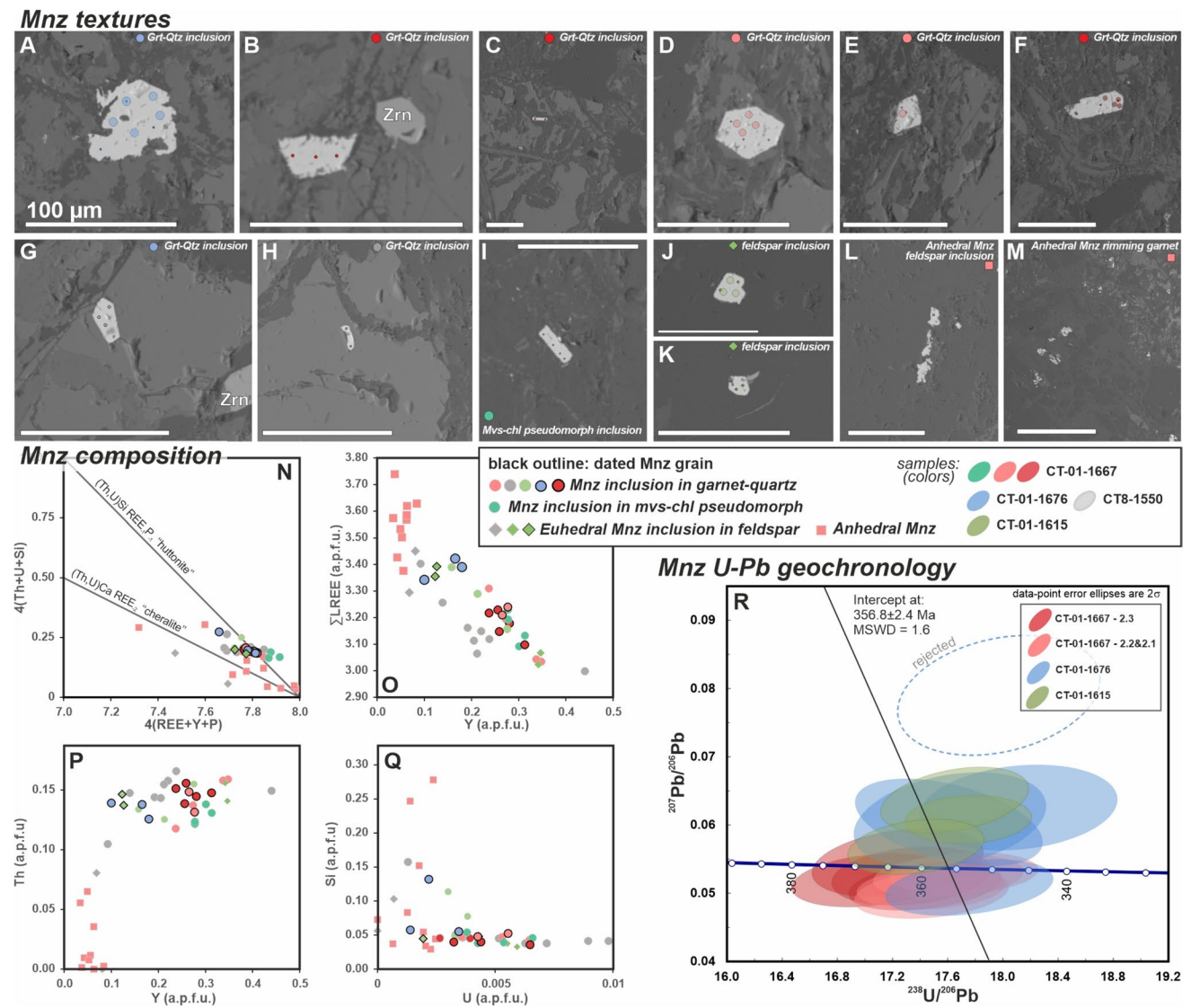


Fig. 13 A–M Monazite BSE images with EPMA and LA-ICP-MS spot analyses. Scale bars are 100 μm ; N–Q EPMA monazite chemical composition; R Tera–Wasserburg concordia diagram of the LA-ICP-MS U–Pb monazite results

garnet intergrown with quartz and one Th-rich euhedral monazite inclusion in feldspar (Fig. 13A, D, E, F, J; ESM2 Table 9; methods in ESM4). One highly discordant analysis ($> 10\%$) was rejected from the dataset (Fig. 13A, R). The remaining sixteen analyses are below 10% discordance, suggesting low common Pb contents. The $^{206}\text{Pb}/^{238}\text{U}$ data define a Tera–Wasserburg lower intercept age of 356.8 ± 2.4 Ma (MSWD = 1.6; Fig. 13R), anchored at 0.859 ± 0.02 , derived from the Stacey and Kramers (1975) terrestrial Pb evolution model. This date overlaps, within uncertainty with the youngest zircon dates from the *ca.* 365 Ma major population of the SHRIMP dataset, and significantly overlaps with the *ca.* 360 Ma age main zircon population in the LA-ICP-MS dataset (Fig. 11).

Geothermobarometric constraints

Ti-in-zircon and zircon saturation temperatures

The reduced zircon $f\text{O}_2$ estimates preclude rutile crystallization ($a\text{TiO}_2 = 1$), which is a typical feature of oxidized melts (e.g., Schiller and Finger 2019). This strongly suggests that $a\text{TiO}_2$ should be below 1. Independent $a\text{TiO}_2$ estimates are not available, especially due to the widespread alteration of primary oxide phases into rutile + titanite + chlorite, possibly resulting from the low-grade metamorphic recrystallization of ilmenite (e.g., Luvizotto et al. 2009; Angiboust and Harlov 2017). Such sub-solidus alteration is also supported by the widespread

porous feldspar textures, and replacement of euhedral albite feldspars by epidote and muscovite (e.g., Plümper and Putnis 2009), hindering the use of oxide phases to estimate magmatic $a\text{TiO}_2$ values. Ilmenite-stable silicic melts have variable $a\text{TiO}_2$, ranging from at least 0.3 up to 0.9 (Teixeira et al. 2024), although typically considered between 0.4 and 0.5 (Schiller and Finger 2019). Furthermore, variations in $a\text{TiO}_2$ are expected during the cooling and evolution of felsic melts, and thus the use of a fixed $a\text{TiO}_2$ may also introduce significant discrepancies in the calculated Ti-in-zircon temperatures (Teixeira et al. 2024). As such, we calculate zircon crystallization temperatures using the Ti-in-zircon calibration of Ferry and Watson (2007), excluding two analysis with anomalously high Ti concentrations (59 and 72 $\mu\text{g/g}$), and considering $a\text{SiO}_2 = 1$ and $a\text{TiO}_2$ at 0.5 and 0.9. Calculated temperatures with $a\text{TiO}_2$ of 0.5 are high and range from 710 up to 965 $^\circ\text{C}$ (Fig. 12F, J), whereas those obtained considering $a\text{TiO}_2$ of 0.9 are lower, and range between ca. 660 and 890 $^\circ\text{C}$. Furthermore, the average temperatures of the ca. 365 Ma population is $837 \pm 66^\circ\text{C}$ ($a\text{TiO}_2$ of 0.5) and $774 \pm 60^\circ\text{C}$ ($a\text{TiO}_2$ of 0.9) (Fig. 12J). These latter estimates ($a\text{TiO}_2$ of 0.9) are thus preferred but are considered as minimum zircon crystallization temperatures (T_{Zirc}).

Experimental work on Zr solubility in felsic melts resulted in the development of zircon saturation models, which enables predicting the temperature at which a melt with a given composition (M factor) saturates in zircon (Watson and Harrison 1983; Boehnke et al. 2013). Most garnet-bearing rhyolites are within the calibrated compositional

range of $0.9 < M < 1.9$ ($M = (\text{Na} + \text{K} + 2\text{Ca})/\text{Al-Si}$ in cation proportions), resulting in zircon saturation temperatures (T_{SatZirc}) between 696 and 800 $^\circ\text{C}$, using the Boehnke et al (2013) model. T_{SatZirc} is inversely correlated with SiO_2 and is positively correlated with $\text{FeO}_T + \text{MgO}$ (ESM2 Table 3), and the most silicic compositions (71.5–78.3 SiO_2 wt%) display T_{SatZirc} between 696 and 772 $^\circ\text{C}$ (ESM2 Table 3). The presence of inherited zircon implies that the melts were zircon oversaturated or near saturation. As such, the T_{SatZirc} are considered as maximum temperatures of the melts. T_{SatZirc} values overlap with most of the minimum zircon crystallization temperatures (T_{Zirc}).

TitaniQ: Ti-in-quartz

Quartz intergrown with garnet shows a bright and homogeneous CL response and exhibits rounded and smooth-walled embayments (Fig. 14A–D). Five LA-ICP-MS analyses of quartz intergrown with garnet show Ti contents between 92 and 108 $\mu\text{g/g}$ (Fig. 14A–D; ESM2 Table 11). Ti incorporation into the quartz lattice is sensitive to temperature, pressure, melt composition and $a\text{TiO}_2$ (e.g., Wark and Watson 2006; Osborne et al. 2022). We use the Ti-in-quartz calibration of Osborne et al. (2022) to model the effects of varying P–T– $a\text{TiO}_2$ conditions (Fig. 14E). Quartz crystallization in equilibrium with rutile ($a\text{TiO}_2 = 1$) is only shown for comparison, as zircon oxybarometry indicates reduced magmatic $f\text{O}_2$ conditions. The average minimum Ti-in-zircon temperature estimates ($a\text{TiO}_2 = 0.9$) are typically lower than those required for dehydration

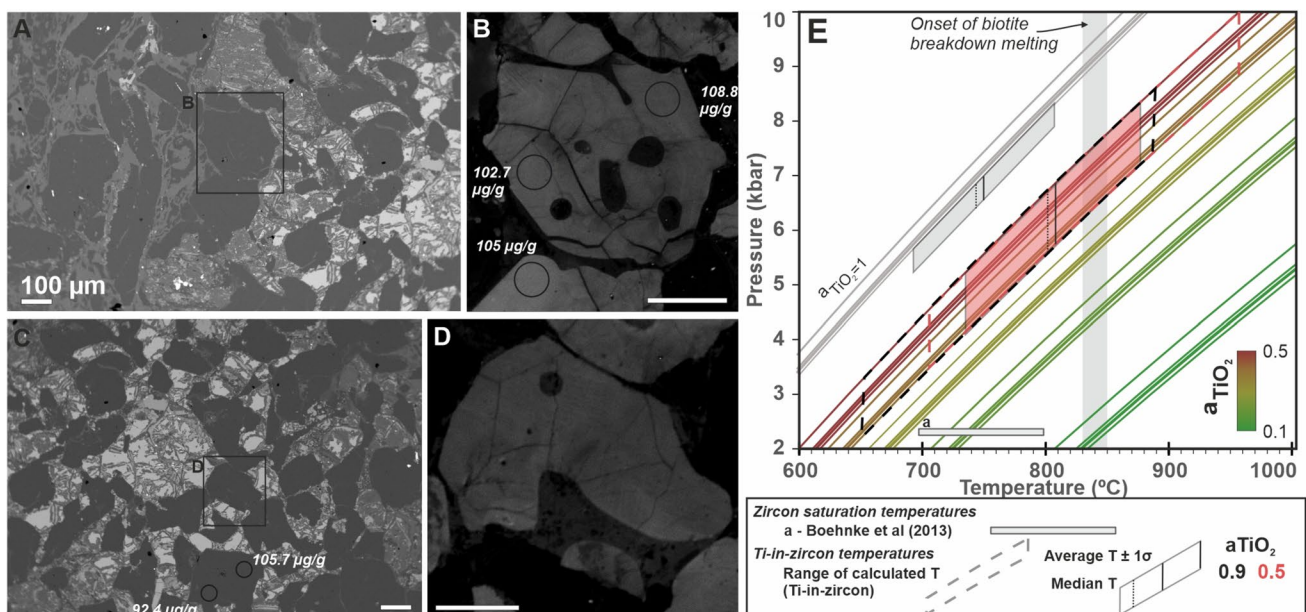


Fig. 14 A–D BSE and cathodoluminescence (CL) features of the analyzed quartz intergrown with garnet; E P-T- $a\text{TiO}_2$ results of the analyzed quartz intergrown with garnet

melting involving biotite (> 820–850 °C), although the highest calculated T_{Zirc} for the 350–370 Ma zircon analyses are compatible with dehydration melting involving biotite (up to 870 °C). Considering these high-temperatures and high aTiO₂ values (0.9), quartz crystallization is calculated at high pressure conditions (8–10 kbar), whereas a lower aTiO₂ of between 0.3 and 0.5 during quartz crystallization results in lower pressure estimates, between 6 and 7 kbar (Fig. 14E). This large range in potential pressure conditions further highlights the importance of constraining aTiO₂. The amount of TiO₂ in a melt (aTiO₂), however, is not constant in natural systems (Schiller and Finger 2019; Teixeira et al. 2024). aTiO₂ in felsic melts may either increase during cooling and crystallization of Ti-poor minerals or decrease due to crystallization and efficient removal of Ti-rich phases. As such, it is likely that different aTiO₂ values should be considered when using the Ti-in-quartz and Ti-in-zircon geothermometry (Teixeira et al. 2024).

Discussion

Garnets in silicic volcanic rocks: origin and growth

Origin of the garnets

While more common in low- to high-grade metamorphic rocks, garnet also occasionally occurs as a rock-forming phase in felsic volcanic (Harangi et al. 2002; Lucci et al. 2018; Sieck et al. 2019) and plutonic rocks (Dahlquist et al. 2007; Villaros et al. 2009a; Dorais and Tubrett 2012; Dorais and Spencer 2014; Narduzzi et al. 2017), and in highly differentiated and pegmatitic felsic rocks (Zhang et al. 2001; Breiter et al. 2005; Samadi et al. 2014). Interpretations on the origin of garnets in felsic volcanic and plutonic rocks range from: (i) xenocrystic garnets, derived from the incomplete assimilation of the protolith or other metamorphic country rocks (e.g., Clarke 2007; Scallion et al. 2011); (ii) a peritectic origin, generated during incongruent partial melting reactions and entrained into the extracted melt (Erdmann et al. 2009; Villaros et al. 2009a; Taylor and Stevens 2010; Dorais and Tubrett 2012; Jung et al. 2022); (iii) a phenocryst/liquidus phase, directly crystallizing from the felsic melt (e.g., Lucci et al. 2018; Sieck et al. 2019; Devoir et al. 2021), including in highly fractionated felsic melts or pegmatites (e.g., Samadi et al. 2014; Maner et al. 2019). Textural, major- and trace element data of these phases provide critical insights into the petrogenesis of the host rocks (Clarke 2007; Erdmann et al. 2009; Dorais and Campbell 2022).

Comparing the major element chemistry from garnets of this study with those compiled in Schönig et al. (2021), the garnets from this study overlap with compositions of

those from felsic igneous rocks and metamorphic rocks in metasediments (ESM3), which are characterized by low Ca and Mn contents (ESM3). Furthermore, metamorphic garnets are commonly characterized by a variety of metamorphic mineral inclusions, usually display textural evidence of deformation and/or metamorphic mineral growth (e.g., snowball structures), and significant major element core-to-rim compositional zoning (e.g., Groppo et al. 2012). Xenocrystic garnets in felsic plutonic and volcanic rocks also frequently show an association with partially disaggregated and assimilated crustal material (Scallion et al. 2011) or show textural and chemical features that indicate significant resorption, replacement or chemical exchange with felsic melts (Clarke 2007; Devoir et al. 2021). The absence of metamorphic mineral inclusions and of pronounced major element core-to-rim zonation and the lack of other xenolithic material strongly argues an origin of the garnets related to assimilated metamorphic country rocks.

Garnets that crystallize from evolved felsic melts sometimes exhibit similar textural features such as quartz intergrowths and rounded- or lobate-shaped inclusions and graphic-like intergrowths (e.g., Zhang et al. 2001; Breiter et al. 2005; Samadi et al. 2014). Despite the textural similarities, these garnets display strong enrichments in Mn and typically X_{Sps} dominates over other components. The role of Mn in stabilizing garnet at lower pressures has been widely documented, enabling magmatic and metamorphic crystallization at low pressures (< 5 kbar) in the upper crust (e.g., Green 1977; White et al. 2014b). The overall low X_{Sps} contents ($X_{\text{Sps}} < 4\%$) of the garnets strongly argues against garnet crystallization during emplacement and melt evolution in a shallow crustal magmatic reservoir (Lucci et al. 2018; Sieck et al. 2019; Devoir et al. 2021), consistent with our Ti-in-quartz pressure constraints that indicate crystallization between 6 and 7 kbar, in the middle–lower crust (Fig. 14).

The textural features of the garnet–quartz intergrowths strongly suggests crystallization in the presence of melt. The abundant quartz inclusions and intergrowths and amalgamated garnet–quartz grains closely resemble those found in granulite-facies migmatites (White et al. 2004; Taylor and Stevens 2010; Taylor et al. 2010; Ferrero et al. 2012; Yakymchuk et al. 2015; Blereau et al. 2017; Goncalves et al. 2021), and also quartz–garnet intergrowths formed during fluid-absent partial melting experiments of biotite-bearing metapelites (Vielzeuf and Montel 1994; Waters 2001; Barbey 2007). Similar peritectic phases and quartz intergrowths can also be found in amphibolite-facies anatectic terranes (Dyck et al. 2019), where fluid-absent muscovite breakdown reactions form peritectic K-feldspar and sillimanite. Film-like, rounded to lobate quartz inclusions and intergrowths are common in peritectic phases, generated during sluggish, kinetically

controlled partial melting reactions under disequilibrium conditions (Waters 2001; Barbey 2007). Quartz intergrown with garnet partially exhibits optical continuity within the garnet–quartz aggregates, but the variable shapes of the quartz intergrowths and inclusions (from film-like and lobate to euhedral inclusions) may also suggest variable effects of post-crystallization shape change and maturation processes (Cesare et al. 2021). However, the CL images of quartz inclusions in garnet show the presence of disequilibrium textural features such as embayments, consistent with rapid quartz crystallization during peritectic garnet growth (e.g., Barbee et al. 2020). The above observations show the garnets are unambiguously of peritectic origin, entrained during melt extraction from the partial melting region.

Despite ample evidence that garnet does not typically survive melt extraction and upper-crustal emplacement and/or eruption due to rapid dissolution–reprecipitation and/or compositional homogenization (Clemens et al. 1997; Villaros et al. 2009a; Taylor and Stevens 2010; Devoir et al. 2021; Jung et al. 2022), our results suggest that diffusion was responsible for major element homogenization of garnet, instead of dissolution–reprecipitation processes. This is further supported by the lack of textural and chemical evidences of dissolution–reprecipitation processes, indicating that peritectic garnets preserve their chemical and textural features. This evidence suggests little interaction and/or re-equilibration of the peritectic garnets with the melt. This may be due to equilibrium with the melt or sluggish diffusivities at the garnet–melt interface, inhibiting dissolution–reprecipitation reactions (Acosta-Vigil et al. 2017), or that the peritectic phases and silicic melt were rapidly extracted from the source region. Indeed, rapid melt loss from the source is required to preserve the anhydrous high-temperature peritectic phases, as melt crystallization and cooling would result in significant garnet back-reaction and replacement (White and Powell 2002; Clemens and Stevens 2016). These processes require the existence of an interconnected melt network in the protolith, which may form at low melt fractions (below 7%; Clemens and Stevens 2016) and possibly enhanced by deformation, ensuring that during the melting reactions the entrained peritectic products are preserved, and transported along with the generated melt.

Trace element zoning and peritectic garnet growth

The trace element LA-ICP-MS spot analyses profiles from garnet–quartz intergrowths are broadly consistent with the zoning pattern shown in the mapped garnet grain. However, the smaller (sub)euhedral garnet grains display somewhat distinct trace element distributions, showing an HREE and Y-poor core (Fig. 8), in contrast to the larger garnets intergrown with quartz (Fig. 7). While it can be argued that the garnet transects do not cross the geometric garnet core,

the major ($X_{\text{Grs}} > X_{\text{Spss}}$) and overall lower trace element concentrations of the smaller euhedral garnets suggest these crystallized after the garnet intergrown with quartz. Hence, we interpret the crystallization of the euhedral garnets during partial melting reactions, after the crystallization of the early peritectic garnet–quartz intergrowths.

The oscillatory zoning displayed in the core and mantle mapped garnet intergrown with quartz could have developed during subsolidus metamorphic garnet growth (Konrad-Schmolke et al. 2022). However, there is no textural and/or compositional evidence for the presence of dissolved xenocrystic sub-solidus garnet cores (e.g., Rubatto et al. 2020; Devoir et al. 2021; Jung et al. 2022). The highest estimated minimum zircon crystallization temperatures (> 800 °C) and the presence of Th–Y-rich monazite inclusions in the core, mantle and rim of the peritectic garnet further shows garnet crystallization at high-temperatures and in the presence of a melt and thus under supra-solidus conditions. The pronounced oscillatory zoning in the garnet core, mantle, and rim in most trace element maps is thus interpreted as resulting from the competing effects of trace element incorporation during peritectic garnet crystallization and the supply rate from the melt via diffusion during partial melting. More importantly, the inner core and mantle are similarly enriched in V, Sc, Zr, Ti and HREE and display sharp compositional gradients. Therefore, the mapped zoning should reflect the breakdown of major and accessory phases during partial melting reactions. While V and Sc indicate the breakdown of biotite during melting reactions (Yang and Rivers 2000; Dorais and Tubrett 2012), the HREE, Zr, Ti and Y zoning attest to the dissolution of accessory phases such as zircon, monazite or apatite during partial melting reactions involving biotite (Dorais and Tubrett 2012; Yakymchuk 2017; Rubatto et al. 2020; Johnson et al. 2021).

Garnet-bearing rhyolites: geochemistry, zircon cargo, and source constraints

Geochemical variation of the garnet-bearing rhyolites

There is a clear lack of magma mixing indicators, such as microgranular intermediate-mafic igneous enclaves, resorption, and disequilibrium textures in feldspar and other minerals, besides garnet and pinitized cordierite. The low Mg#, Zr contents and other geochemical features (e.g., the strongly peraluminous character) also attest to magma mixing processes with mafic or intermediate melts as not being a significant process in the generation of the IPB garnet-bearing rhyolites (e.g., Mitjavilla et al. 1997).

The broad positive correlation of K_2O vs $FeO_T + MgO$ is an unusual feature among granitoids and S-type volcanic rocks (Clemens and Stevens 2012) and may indicate a role of

biotite fractionation (Clemens et al. 2017a). However, biotite is scarce in all the samples, and its modal amount is not compatible with significant amounts of biotite fractionation. The positive correlation of K and maficity and the constant K/Rb suggests no fractionation of K-feldspar and overall low K melts, whereas the increase of peraluminosity and Yb with maficity highlights the absence of garnet, cordierite and biotite fractionation (e.g., Champion and Bultitude 2013), in agreement with the fact that garnet and cordierite are peritectic and not liquidus phases. Considering all the plots in Fig. 9, no coherent fractionating assemblage seems capable of reproducing the main major element geochemical trends, and thus, fractional crystallization is ruled out as a main driver of the geochemical variation of the garnet-bearing volcanic rocks.

Recently, several studies have proposed and shown that the geochemical variability of S- and I-type volcanic and plutonic rocks reflects, in some cases, the entrainment of peritectic phases (Stevens et al. 2007; Villaros et al. 2009b; Dorais and Spencer 2014; Clemens et al. 2017b; Bailie et al. 2020; Zhu et al. 2021; Dorais and Campbell 2022). The range in maficity and the positive correlations with A/CNK, TiO₂, Y, Yb, Sm, Zr, Hf, Th, La, Eu and Ce are all hallmarks of PAE. More specifically, the negative correlation of Ca with maficity suggests entrainment of Ca-poor garnet (Stevens et al. 2007), which is similar to the composition of the entrained peritectic garnets in this study. Furthermore, the positive correlation between maficity and A/CNK, K and Y, Yb, Sm and Nd suggests the entrainment of peritectic cordierite, K-feldspar and garnet. The small slope in the Ti vs FeO_T + MgO plot (Fig. 9F) further indicates a small proportion of peritectic ilmenite entrainment, also in accordance with the textural and mineralogical data. Furthermore, the negative anomalies in K and Rb further support the presence of peritectic/residual K-feldspar. The lack of knowledge on the chemical composition of the protolith, the melt it generated and the composition of the peritectic phases other than garnet, means that the modeling presented in Fig. 9 can only provide a semi-quantitative/qualitative approach of the role of PAE. Nevertheless, the geochemical trends defined by the garnet-bearing rhyolites can be satisfactorily reproduced by entrainment of up to 15% of peritectic garnet, cordierite, and K-feldspar, and less than 1% peritectic ilmenite, into a highly silicic melt. The systematic positive correlation of Zr, Hf, La, Ce and zircon saturation temperatures with maficity also highlights the role of zircon and monazite co-entrainment, generated and/or liberated from the breakdown of biotite during melting reactions (e.g., Villaros et al. 2009b), which are common inclusions in the entrained peritectic garnets and pinitized cordierite and also in intergrown quartz and in feldspar crystals. Ilmenite and Ti-rich biotite are the most common residual phases generated during partial melting of

metasedimentary rocks (White et al. 2014a) and are often found in high-temperature migmatites. The strongly negative Ti anomalies compared to continental crust, together with the marked Ta and Nb (Ta < Nb) anomalies suggest that ilmenite and Ti-rich biotite were residual phases during melting reactions and were not significantly entrained into the extracted melt (Stepanov et al. 2014). The geochemical trends defined by the garnet-bearing rhyolites thus mainly reflect entrainment of peritectic minerals (garnet, cordierite, K-feldspar and minor ilmenite), generated during partial melting reactions involving fluid-absent breakdown of biotite into a highly silicic melt.

Protracted zircon cargo: source area, incremental pluton growth and geochemistry

The zircon U–Pb data from the garnet-bearing rhyolites show a wide range of apparent ages, a common feature of crustal-derived granitic magmas (Villaros et al. 2012; Ferreira et al. 2019; Clemens et al. 2023; Vogt et al. 2023). Furthermore, the U–Pb dates in both the LA-ICP-MS and SHRIMP datasets show significant scatter but define overlapping zircon populations, and are interpreted by integrating the available geological (geochronological, palynological and lithostratigraphic) data.

The youngest zircon population (ca. 345 Ma, only seen in the larger LA-ICP-MS dataset) constrains the eruption age of the felsic volcanic rocks, and agrees with the available palynological and stratigraphic data (mid and upper VSC sediments) estimated at ca. 350–330 Ma (Figs. 3, 11; Oliveira et al. 2004; Carvalho et al. 2017). The main zircon population at ca. 365 Ma (which encompass zircon grains with ages from ca. 350 to 370 Ma) overlaps with the age of a major explosive volcanic episode in the Neves-Corvo area (rhyolite fiamme-facies volcanic rocks; Rosa et al. 2008), interbedded with the Corvo formation, dated from ca. 363 to 365 Ma (Oliveira et al. 2004; Pereira et al. 2021). These analyses might also include younger zircon grains which are not analytically distinguishable from the ca. 365 Ma major population due to the individual uncertainties (e.g., Large et al. 2020) dataset. This is further supported by the varying Y/Hf and Y/Ti trends and Gd/Yb ratios of the zircon analyses, some of which display flat REE patterns, suggesting HREE partition into garnet prior to zircon crystallization (e.g., Taylor et al. 2015). The zircon geochemistry dataset also shows that Y is positively correlated with the Gd/Yb ratios, further supporting the role of garnet crystallization in controlling Y-HREE contents of zircon. Additionally, the high Th of the analyzed monazite grains (see discussion below), and positive correlation of zircon Ti with Th/U also suggests that monazite crystallization controlled the Th/U ratios of magmatic zircon (Pineda et al. 2022; Vogt et al. 2023).

In both datasets, the individual zircon populations overlap, within uncertainty, with published zircon U–Pb ages in the Neves-Corvo and Rosário anticline, and also in other sectors of the IPB (Rosa et al. 2009; Albardeiro et al. 2023). The zircon populations inherited from the melt source region (400–414 Ma; 420–440 Ma; 450–470 Ma) overlap within uncertainty with published zircon U–Pb ages in the Neves-Corvo and Rosário anticline, and also in other sectors of the IPB (Barrie et al. 2002; Rosa et al. 2009; Valenzuela et al. 2011; Lains Amaral et al. 2021; Pereira et al. 2021; de Mello et al. 2022; Albardeiro et al. 2023). The inherited zircon dates also broadly overlap with zircon ages found throughout Avalonia and in the Meguma terranes, especially with Ordovician metasediments from the upper Meguma Supergroup (Murphy and Hamilton 2000; Waldron et al. 2009; Braid et al. 2012; Shellnutt et al. 2019), further supporting a relationship between the peri-Gondwana Meguma terrane and the unexposed SPZ basement (e.g., Braid et al. 2012).

Palynological constraints from the oldest known PQG metasediments places the onset of sedimentation in the IPB to the Middle Devonian (387–384 Ma) (Mendes et al. 2020), which indicates that zircon crystals younger than *ca.* 390 Ma cannot be source-inherited or derived from other country rocks. Instead, the abundant presence of pre-eruption Devonian zircon grains can only be explained by the remobilization and extrusion of previously crystallized felsic melts (e.g., Albardeiro et al. 2023). These younger zircon populations overlap remarkably with major volcanic episodes in the Rosário–Neves-Corvo anticline and in other sectors of the IPB (Fig. 11), namely: *ca.* 360 and 365 Ma in the Neves–Corvo–Rosário region (Rosa et al. 2008; Pereira et al. 2021), and *ca.* 370–376 in the Cercal region (Rosa et al. 2009). Older zircon ages (*ca.* 390–380 Ma) also overlap with previously recognized “xenocrystic” U–Pb zircon ages in felsic volcanic rocks of the IPB (Barrie et al. 2002; Rosa et al. 2009; Valenzuela et al. 2011; Lains Amaral et al. 2021; Pereira et al. 2021; Albardeiro et al. 2023). These medium- to high-temperature reduced felsic melts were thus stored at temperatures below the solidus in an upper-crustal pluton which records the episodic emplacement and crystallization of felsic melts in the upper crust since *ca.* 390 Ma, possibly marking the onset of the intracontinental rifting in the South Portuguese Zone basement.

Evidence for high-temperature partial melting of the middle crust

Peritectic assemblages formed after biotite breakdown reactions are dominated by garnet, cordierite and/or orthopyroxene, depending mainly on the range of pressures and protolith compositions (e.g., Johnson et al. 2021).

Therefore, the entrained peritectic garnet and cordierite indicate incongruent melting reactions involving biotite.

While the low X_{Pyr} contents of the entrained peritectic garnets may suggest low-temperature conditions, as Mg in garnet increases with temperature due to the Fe–Mg exchange with biotite, Fe-rich garnet compositions are common in the middle crust, due to Fe–Mg partitioning with coexisting cordierite (Hensen and Green 1972; Holdaway and Lee 1977). Cordierite is a common phase in low-pressure (< 5 kbar) metasedimentary migmatites and S-type granites (Clarke 1995; Barbey et al. 1999; Groppo et al. 2013; White et al. 2014b), but the stability of (Mg-rich) cordierite extends up to 7–8 kbar, in equilibrium with garnet, following the melt-producing reaction: biotite + aluminosilicate + quartz + albite = garnet + cordierite + K-feldspar + melt (Holdaway and Lee 1977; Spear et al. 1999; Vielzeuf and Schmidt 2001), and its stability should be expanded by increasing protolith Mg#. Hence, we interpret the low Mg- and high Fe as resulting from partial melting at temperature conditions near the onset of biotite dehydration melting (e.g., Ward et al. 2008) and from peritectic Grt–Crd equilibrium. This is consistent with the presence of pinitized cordierite(?)–quartz intergrowths, which are texturally similar to the peritectic garnet–quartz intergrowths, also indicating its peritectic nature. Furthermore, even though Ca contents in garnet from S-type melts are typically low, peritectic garnets in high-pressure granulite-facies migmatites commonly display higher X_{Grs} with increasing pressure even in cases of partial melting of low Mg# metapelitic protoliths (e.g., Li et al. 2019). This further suggests partial melting at mid-crustal pressures.

The use of the Ti-in-quartz solubility model in estimating the pressures of quartz crystallization during early peritectic garnet crystallization yields widely varying pressures, from 8 to 10 kbar when considering high aTiO₂ (0.9, also used to calculate T_{Zirc}), to 6–7 kbar if lower aTiO₂ (0.5–0.4) are considered. The 8–10 kbar pressure range is significantly higher than that required for cordierite to be stable, and the pressure conditions of 6–7 kbar provide more reasonable estimates. This, however, implies variations in aTiO₂, which is common in evolving silicic melts (Teixeira et al. 2024). As such, considering varying aTiO₂ for quartz and zircon crystallization may provide a more realistic approach to the use of Ti-based geothermometers (Teixeira et al. 2024). Our quartz Ti contents are typically below 200, which suggest crystallization at aTiO₂ lower than 0.8 (Teixeira et al. 2024). The low Ti contents of the felsic volcanic rocks and our PAE modeling further suggests a low aTiO₂ for the melts, possibly due to efficient removal and/or lack of equilibrium of the melt and residual ilmenite, resulting in overall low aTiO₂ during the early partial melting reactions. After melt extraction from the source, however, cooling

and crystallization of other phases would likely result in increasing Ti contents of the melts (Teixeira et al. 2024).

The absence of additional $a\text{TiO}_2$ constraints, preserved minerals in equilibrium with garnet, the pervasive alteration and replacement of cordierite, and lack of whole-rock compositions that resemble the melt from which the phases crystallize hinders the use of garnet-based geothermometers or pseudosections to precisely constrain P–T conditions during partial melting. Nevertheless, the geothermobarometric constraints from zircon and quartz intergrown with garnet are consistent with peritectic garnet and cordierite crystallization during fluid-absent incongruent melting reactions involving biotite of a metasedimentary protolith in the middle crust (6–7 kbar), at temperature conditions near the onset of biotite incongruent melting reactions (*ca.* 820–870 °C).

The major element zoning of the entrained peritectic garnets mimics common growth zonation of peritectic garnets generated during high-temperature incongruent partial melting reactions (Cesare 2000; Groppo et al. 2012). Garnet zoning is mainly controlled by the diffusion rate during growth (Clarke 1981). At temperatures below 640 ± 50 °C, major element diffusion is negligible (Yardley 1977), whereas at temperatures above 700 °C, diffusion rates of Fe, Mn and other major elements exponentially increase. Our Ti-in-zircon (T_{Zirc}) thermometry results indicate magmatic zircon crystallization at high temperatures, between *ca.* 690–870 °C, but the preservation of garnet growth zoning suggests a limited residence time at high-T conditions (> 700 °C), preventing chemical exchange and interaction with the host melt. This further implies relatively fast extraction rates from the partial melting region and migration through the crust, which is possibly the cause of the intense garnet fracturing. This, however, would likely promote reaction with the melt and replacement. Given the lack of cordierite and/or biotite-dominated replacements of garnet along the fractures, it is also possible that garnet fracturing was due to increasing pressure (above lithostatic pressure) conditions, during magma recharge and eruption.

Additionally, the reduced estimated $f\text{O}_2$ magmatic conditions and the aluminous peritectic assemblages further indicate that the melt and peritectic phases were generated through the partial melting of reduced aluminous metasedimentary protoliths. Although melting of metagraywackes can produce garnet and cordierite as peritectic phases, in the range of the previously discussed P–T estimates (< 870 °C and 6–7 kbar) orthopyroxene typically dominates peritectic assemblages (Vielzeuf and Montel 1994), indicating partial melting of a reduced aluminous pelitic source in the middle crust.

Such high-temperature melting processes in the middle crust indicates high geothermal gradients (above 40 °C/

km), a scenario previously suggested for the Paleozoic Iberian Pyrite Belt, a significantly base metal-endowed mining district (Tornos 2006). Such high geothermal gradients are likely induced by the mid to lower crustal emplacement of hot mantle-derived melts (Mitjavilla et al. 1997), providing the required heat to melt the crust, while deformation associated with the transtensional opening of the IPB basins may have further enhanced melt extraction and upper-crustal emplacement (Brown 2013; Clemens and Stevens 2016; Clemens et al. 2020; Johnson et al. 2021).

Preserved suprasolidus prograde monazite

Monazite in felsic magmatic rocks commonly grows during magma cooling and crystallization, after melt extraction from the source (Spear and Pyle 2010; Yakymchuk 2017). Suprasolidus prograde monazite has, nonetheless, been documented in migmatites (*e.g.*, Johnson et al. 2015; Blereau et al. 2016; Rocha et al. 2017).

The higher partitioning of Th into monazite in equilibrium with a melt (Xing et al. 2013) indicates that the high-Th monazite inclusions in the peritectic garnets and pinitized cordierite crystallized from a melt. Although the lack of monazite HREE data from the analyzed monazite grains hinders a complete assessment of the monazite-garnet equilibrium, the Y-rich composition of these inclusions further indicates monazite crystallization prior to peritectic garnet, as Y is highly compatible in garnet (*e.g.*, Pyle and Spear 2003). The Th–Y-rich monazite inclusions in peritectic garnets are thus interpreted as preserved prograde suprasolidus monazite, crystallized during earlier stages of partial melting. As temperature and melt connectivity increases in the partial melting area, prograde monazite is often subsequently dissolved into the melt (Yakymchuk 2017). However, the preservation potential of monazite can increase if included into co- or subsequently precipitated peritectic minerals, such as garnet. This is the mechanism most likely for the preservation of the suite of monazite grains included in garnet and pinitized cordierite. Therefore, the U–Pb monazite age of 356.8 ± 2.4 Ma is interpreted as representing the timing of peritectic garnet growth during partial melting reactions and overlaps within uncertainty with the *ca.* 360 Ma zircon population in the LA-ICP-MS dataset (Fig. 11), which we interpret as the timing of garnet growth. Although increasing evidence shows that the duration of partial melting and felsic melt extraction and emplacement processes are typically fast (< 1 Myr; Coulson et al. 2002; Devoir et al. 2021; Jung et al. 2022), felsic melts may also have long residence times in the lower to upper crust, prior to eruption (*e.g.*, Cesare et al. 2003, 2009) as discussed below.

Eruption and preservation of peritectic phases

Our results clearly show that the erupted melts carried a significant amount of previously crystallized crystal cargo from older melt batches, which also fed voluminous eruptions. This attests to the presence of an upper-crustal reservoir that stored crystallized melts from previous voluminous crustal melt generation, storage and eruption episodes. Their preservation and eruption, however, requires that these phases survive subsequent melt recharge and extrusion, at *ca.* 345 Ma, after *ca.* 15 Ma of storage below the solidus.

The lack of geochemical data from the youngest zircon grains hinders a complete assessment of some critical intensive parameter of the erupted melts such as temperature. Nevertheless, their clear K-poor and silicic nature, combined with the lower zircon saturation temperatures of the most silicic whole-rock compositions (700–770 °C) suggests these melts were generated at lower temperatures and possibly from a biotite-poor source in the middle crust. This points to fluid-absent muscovite breakdown of a lower-grade protolith, which also commonly results in the crystallization of peritectic/residual K-feldspar (e.g., Dyck et al. 2019). Regardless of the nature of the source, the lower temperatures of the erupted melt would likely inhibit significant crystal cargo dissolution, offering a plausible explanation for the preservation of the abundant xenocrysts of older peritectic phases. Such scenario closely resembles the El Hoyazo dacites (Acosta-Vigil et al. 2010), which extruded at *ca.* 6 Ma a suite of remelted older (9 Ma) granulite-facies middle to lower crust glass-rich xenoliths, with abundant peritectic phases such as garnet and cordierite. As discussed in Acosta-Vigil et al (2010), the upper-crustal emplacement of felsic melts and entrained peritectic phases would largely induced melt crystallization, whereby subsequent water exsolution would remove most H₂O from the melt. As such, during subsequent voluminous magma recharge and consequent eruption of lower-temperature felsic magma, at *ca.* 345 Ma, melting, dissolution and replacement of the stored crystallized melts and peritectic phases would be inhibited, further promoting the preservation of the abundant xenoliths up to eruption.

Conclusions

In this study, we integrate geological, petrological, major and trace element whole-rock geochemical data of garnet-bearing rhyolites with major and trace element composition of garnet, Ti-in-quartz intergrown with garnet, trace element analysis and U–Pb isotopic dating of monazite and zircon to reach the following conclusions:

- (1) Textural and chemical data from garnets-quartz intergrowths and smaller euhedral garnets clearly show that they represent entrained peritectic phases. These were generated during incongruent melting reactions involving biotite of a pelitic source in the middle crust, at temperatures up to 870 °C.
- (2) Garnet major element composition show homogenization by diffusion processes, but growth zoning is still preserved in large (up to 1 cm) garnet grains. The overall Fe-rich and Mg-poor composition of the peritectic garnets indicates compositional equilibrium with peritectic cordierite, which is also entrained and preserved as pinitized cordierite–quartz intergrowths.
- (3) Despite the limited major element variation, garnets also exhibit prominent trace element oscillatory zoning and significant core-to-rim variations, recording the breakdown of biotite at high temperatures and also the dissolution of accessory phases during melting reactions.
- (4) The preservation of the peritectic phases and lack of evidence for compositional modification via dissolution–reprecipitation processes suggests rapid melt extraction from the source, but also no significant compositional re-equilibration with the melt during extraction up to extrusion.
- (5) The Th- and Y-rich monazite inclusions in peritectic garnet and cordierite are interpreted as preserved prograde suprasolidus monazite and place the timing of partial melting at 356.8 ± 2.4 Ma. This age overlaps with the 360 ± 1 Ma younger zircon population and represents one of the older major volcanic episodes in the region.
- (6) Our data shows that the extruded silicic melts also carried a significant amount of pre-eruption zircon cargo, containing not only source-inherited zircon crystals, but also zircon that crystallized from previous moderate to high-temperature reduced felsic melts, possibly stored in an episodically and incrementally grown pluton, since *ca.* 390 Ma.
- (7) The whole-rock geochemical data of the garnet-bearing volcanic rocks mainly reflects the entrainment of peritectic phases such as garnet, cordierite and K-feldspar, and our results thus show that a significant amount of xenoliths of peritectic and accessory phases from previously crystallized felsic melts were extruded along with a lower-temperature silicic magma, at *ca.* 345 Ma.
- (8) This work further supports the view of protracted and incremental growth processes of magmatic reservoirs by accretion of several melt batches in a subsolidus low-temperature reservoir, which may be partly remo-

bilized and extruded during subsequent melt recharge and extrusion episodes.

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Data availability The data of this study are available in the Electronic Supplementary Materials.

Declarations

Conflict of interest The authors declare that they have no conflicts of interest.

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