



Archaean phosphates: a case study of transformation processes in apatite from the Barberton greenstone belt

Łukasz Birski¹ · E. Słaby¹ · R. Wirth² · M. Koch-Müller² · K. Simon³ · A. Wudarska^{1,2} · J. Götze⁴ · A. Lepland^{5,6,7} · A. Hofmann⁸ · A. Kuras²

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Abstract

Multiple tools have been used to determine the sensitivity of phosphates from the early Archaean Barberton greenstone belt to transformation. The assessment of the degree of transformation is crucial for verifying data about the parameters of the paleo-environment. From the obtained results, three generations of phosphates can be distinguished. Group A is observed in cherts and banded iron formation BIF early-generation fluor-hydroxyapatite that precipitated from seawater. It is characterized by flat rare earth element (REE) patterns with a positive Eu anomaly and high Y/Ho ratio in the range of 54–70. Apatites in this group lack any visible indicators of secondary alterations at the micro- and nanoscales. Fourier transform infrared spectra indicate that these apatites are relatively rich in water, and, due to cationic substitution, their OH-stretching regions exhibit complex ordering and numerous component bands. The characteristics observed in the cherts and silicified felsic volcanoclastics of group B imply advanced metasomatic alteration. They exhibit light and heavy REE depletion and an absence of water in the halogen site. Nanoscale investigations reveal cracks, pores, nanofluid inclusions and nanochannel-like structures, as well as inclusions. Group C is represented by igneous-derived apatites that partially reflect their igneous origin. The phosphates are predominantly fluorapatite with typical magmatic apatite REE distribution patterns. Imaging at the micro- and nanoscales indicates that they partially preserve the signature of igneous origin. It seems that some of the analyzed apatite partially preserved their primordial features; therefore, they might be used for the reconstruction of Archaean abiotic systems.

Keywords Archaean · Barberton · Apatite · Volatiles · REE · Metasomatism

Introduction

The calcium phosphate mineral apatite, with the general formula $\text{Ca}_{10}(\text{PO}_4)_6(\text{F},\text{Cl},\text{OH})_2$, occurs in a wide range of geologic and biologic settings on Earth. Due to its ubiquity and ability to incorporate water, halogens, S, C, and trace

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✉ Łukasz Birski
l.birski@twarda.pan.pl

¹ Institute of Geological Sciences, Research Centre in Warsaw, Polish Academy of Sciences, Twarda 51/55, 00-818 Warsaw, Poland

² GFZ German Research Centre for Geosciences, Telegrafenberg, Potsdam 14473, Germany

³ University of Göttingen, Goldschmidtstraße 1, 37077 Göttingen, Germany

⁴ TU Bergakademie Freiberg, Institute of Mineralogy, Brennhausgasse 14, 09596 Freiberg, Germany

⁵ Geological Survey of Norway, Leiv Eirikssons vei 39, 7491 Trondheim, Norway

⁶ Department of Geology, University of Tartu, 50411 Tartu, Estonia

⁷ Institute of Geology, Tallinn University of Technology, 19086 Tallinn, Estonia

⁸ Department of Geology, University of Johannesburg, Auckland Park, Johannesburg 2006, South Africa

elements including Sr, U, Th, and the rare earth elements (REEs), apatite is a powerful indicator of primary formation and transformation processes. Phosphate of sedimentary origin precipitates from seawater via organic matter sedimentation and by scavenging onto iron–manganese oxide particles (Filippelli 2008). Apatite of igneous origin crystallizes from melts (Webster and Piccoli 2015), whereas metasomatic apatite is formed due to interaction with fluids (Harlov 2015).

Many papers have studied phosphates of Precambrian age (e.g., Mojzsis et al. 1996; Lepland et al. 2002, 2013; Nishizawa et al. 2004, 2005; Blake et al. 2010; Joosu et al. 2015, 2016; Alibert 2016; Wudarska et al. 2018) with the scope to deduce their primary formation conditions. In the oldest known supracrustal rocks from the Isua Supracrustal Belt (≈ 3.7 Ga) (Moorbath et al. 1973; Nutman et al. 1997), metamorphism (e.g., Rosing et al. 1996; Rollinson 2003) caused obliteration of primordial isotopic signatures (Wudarska et al. 2018), whereas REE signatures have been partially preserved (Lepland et al. 2002; Wudarska et al. 2018). In contrast, the Barberton greenstone belt (BGB) in southern Africa (3.55–3.22 Ga), one of the oldest well-preserved volcanic and sedimentary sequences on Earth, has been metamorphosed to relatively low grades. The carbonaceous material in these rocks has been interpreted as being of biological origin and may contain the oldest recognized microfossils (e.g., Walsh 1992; Walsh and Lowe 1999; Westall et al. 2001; Javaux 2006; Hickman-Lewis et al. 2018). These unique features make the rocks of the BGB ideal candidates to investigate apatite formation conditions during the early Archean. Because apatites/phosphates show some sensitivity to transformation, we try on the basis of their composition and structure to determine the degree of their susceptibility to secondary changes.

The objective of our study is to use a set of compositional and structural data of apatite for the reconstruction of Archean phosphate formation environments, including the behavior of volatiles. It is an attempt to analyze and describe features that are characteristic of primary phosphates of sedimentary and igneous origin not influenced by secondary processes and of phosphates that were metasomatically transformed, e.g., overprinted by interaction with fluids. Most of the samples in our research were previously analyzed for $\delta^{18}\text{O}$ phosphate by Blake et al. (2010). Special emphasis was placed on the volatile compositions and concentrations as their positions in the crystal structures (electron probe microanalyzer—EPMA, Fourier transform infrared spectroscopy—FTIR). The structure and its transformation were investigated by transmission electron microscopy (TEM), and an auxiliary tool was cathodoluminescence (CL). Paleoproterozoic apatite from the Zaonega formation is used as a reference material, giving insight into the relationship between the primary/secondary signatures

resulting from the formation/transformation of the crystals (Joosu et al. 2015, 2016).

Geological setting

The BGB is located in the Kapvaal Craton of southern Africa (Fig. 1), one of the oldest pieces of the continental crust. The BGB consists of a NE–SW-striking succession of Archean supracrustal rocks, termed the Swaziland Supergroup, which is surrounded and intruded by granitoids (Viljoen and Viljoen 1969; de Ronde and de Wit 1994; Lowe and Byerly 2007). The Swaziland Supergroup is subdivided into three stratigraphic units, the Onverwacht, Fig Tree and Moodies groups (Fig. 2). These rock units were tightly folded into a number of synclines that are separated by narrow anticlines or shear zones. The metamorphic grade is greenschist facies but locally reaches amphibolite facies close to contact with the surrounding granitoid domes that intruded during several episodes of magmatism at 3.45, 3.22, and 3.1 Ga (Kamo and Davis 1994; Dziggel et al. 2002). The Onverwacht Group formed between 3.55 and 3.26 Ga (Kröner et al. 2016; Byerly et al. 2018) and consists of ultramafic and mafic volcanic rocks (komatiites, komatiitic basalts, and basalts), with minor dacitic volcanic and sedimentary rocks that formed in a deep-to-shallow-marine environment (Fig. 2). The Onverwacht Group is subdivided into the Sandspruit, Theespruit, Komati, Hooggenoeg, Kromberg and Mendon formations (Fig. 1) (Viljoen and Viljoen 1969; Lowe and Byerly 2007). The Sandspruit and Theespruit Formations are separated from the overlying units by a major fault zone, but the younger units are stratigraphically conformable. Thrust faults that were active during and following the deposition of the Moodies Group are common in the uppermost sedimentary package. The Fig Tree Group, 3.26–3.23 Ga, consists of a several kilometer thick siliciclastic sedimentary sequence that is capped by felsic volcanic rocks (Hofmann 2005). In the northern part of the belt, the Fig Tree Group is characterized by turbiditic sandstone and shale interbeds that formed in a relatively deep-water environment (Hofmann 2005). In the southern part of the belt, a variety of lithofacies, including shale, graywacke, banded iron formation (BIF), conglomerate, and barite, occur that formed in a deep-to-shallow-water to alluvial environment (Fig. 2). The Moodies Group was probably deposited close to 3.227 Ga but not later than 3.11 Ga (Kamo and Davis 1994) and consists of shallow-marine to fluvial sandstone and conglomerate with minor shale and BIF.

In this study, we analyzed some of the samples of Blake et al. (2010) from sedimentary and felsic volcanoclastic units of the Kromberg and Mendon Formations and the Fig Tree Group as well as two additional samples of metachert and dacite-derived sandstone from the Hooggenoeg Formation.

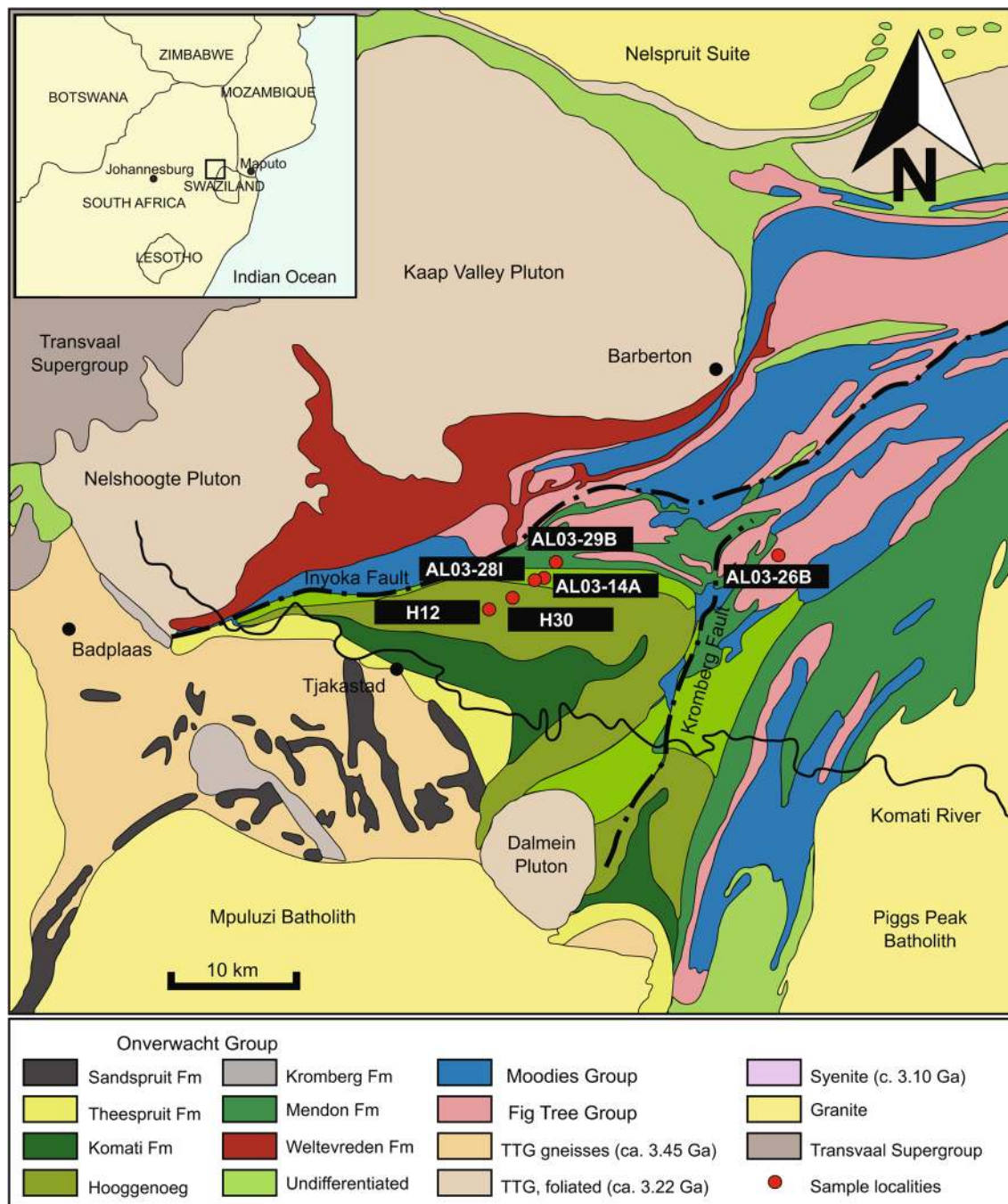


Fig. 1 General geological map of the southwestern part of the Barberton greenstone belt (BGB), South Africa and Swaziland, adopted from Blake et al. (2010). The red dots indicate the sample locations

Sample description

The main idea of our sampling was to reflect the character of the sedimentary and felsic units of the BGB and their petrologic and geochemical characteristics. Therefore, the analyzed samples cover all the complexity and diversity of the upper part of the BGB (Table 1; Figs. 1, 2). We selected six samples from units within the Hooggenoeg, Kromberg

and Mendon formations of the Onverwacht Group and the Fig Tree Group (Fig. 2) that were visually the most representative, free of secondary alterations and therefore able to preserve the primordial signature of the environment of their origin.

Sample AL03-28I was collected from the Footbridge Chert in the Kromberg Formation (Blake et al. 2010) and is a carbonaceous metachert. Quartz is the major phase in

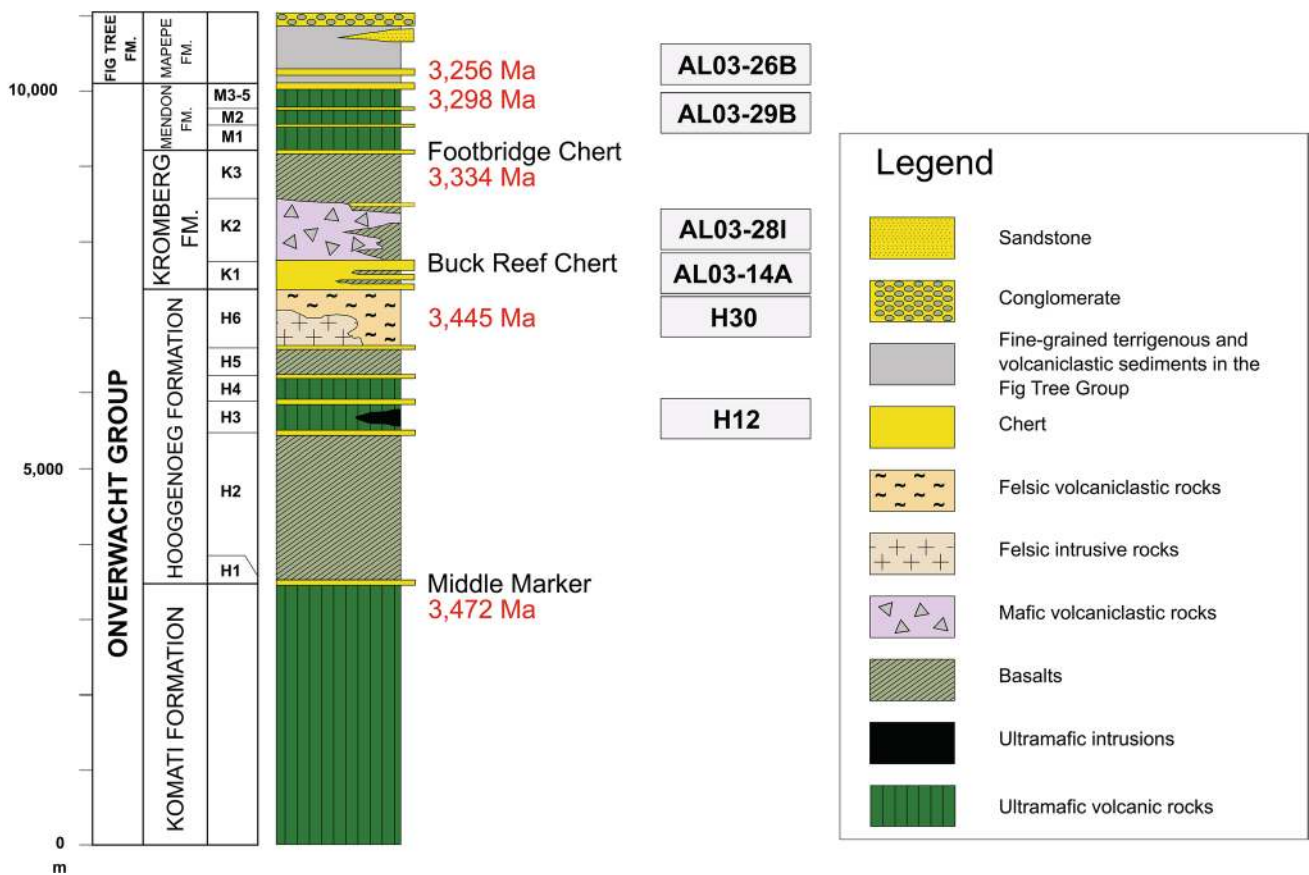


Fig. 2 Generalized stratigraphy and geochronology of the rocks in the upper part of the Onverwacht Group and Fig Tree Group in the southern part of BGB, modified after Knauth and Lowe (2003)

Table 1 Geographical coordinates, lithology, stratigraphy, and mineral composition of the studied samples

Name	Latitude N	Longitude W	Lithology	Stratigraphy	Major phases (> 5%)	Minor phases (< 5%)
AL03-28I	25°55.63	30°54.99	Banded ferruginous chert	Kromberg Formation	Quartz	Haematite, magnetite, calcite, dolomite, pyrite, apatite
AL03-14A	25°55.69	30°54.72	Carbonaceous chert	Kromberg Formation	Quartz	Haematite, magnetite, calcite, dolomite, graphite, pyrite, apatite
H12	25°56.68	30°53.40	Black banded chert	Hooggenoeg Formation	Quartz	Siderite, haematite, calcite, chlorite, apatite, dolomite, rhodochrosite, pyrite
AL03-29B	25°54.86	30°55.85	Silicified felsic volcanoclastics	Mendon Formation	Quartz	Chlorite, apatite, magnetite, pyrite, siderite, dolomite, rhodochrosite, calcite, titanite, zircon
H30	25°55.50	30°53.65	Dacite-derived, silicified sandstone	Hooggenoeg Formation	Quartz, siderite	Sericite, haematite, apatite, dolomite, rhodochrosite, calcite, pyrite, titanite, monazite, zircon
AL03-26B	25°54.12	31°06.35	BIF	Fig Tree Group	Quartz, haematite, magnetite	Calcite, dolomite, pyrite, apatite, rhodochrosite

this sample, and the minor phases are represented mostly by Fe-oxides, i.e., hematite and magnetite (Fig. 3). Locally, in iron-enriched layers, Fe-oxides became the major phase. In addition, calcite, dolomite and pyrite are present. Apatites are very rare, highly porous and small (<30 μm) in the analyzed sample (Table 1; Fig. 3).

Sample AL03-14A is carbonaceous metachert from the Buck Reef Chert in the Kromberg Formation (Blake et al. 2010). It is predominantly composed of quartz. Accessory minerals are represented by Fe-oxides (hematite and

magnetite), carbonates (calcite and dolomite) and pyrite (Fig. 3). Furthermore, the sample is characterized by a relatively high concentration of organic matter and/or organic carbon (Blake et al. 2010). The apatites observed in sample AL03-14A are also very scarce and small (<30 μm) (Table 1; Fig. 3).

Sample H12 is a black-banded metachert from the Hooggenoeg formation (Hofmann et al. 2013). Quartz is the major phase in this sample. Hematite, together with some magnetite, is the most common Fe-oxide (Fig. 3). Rare

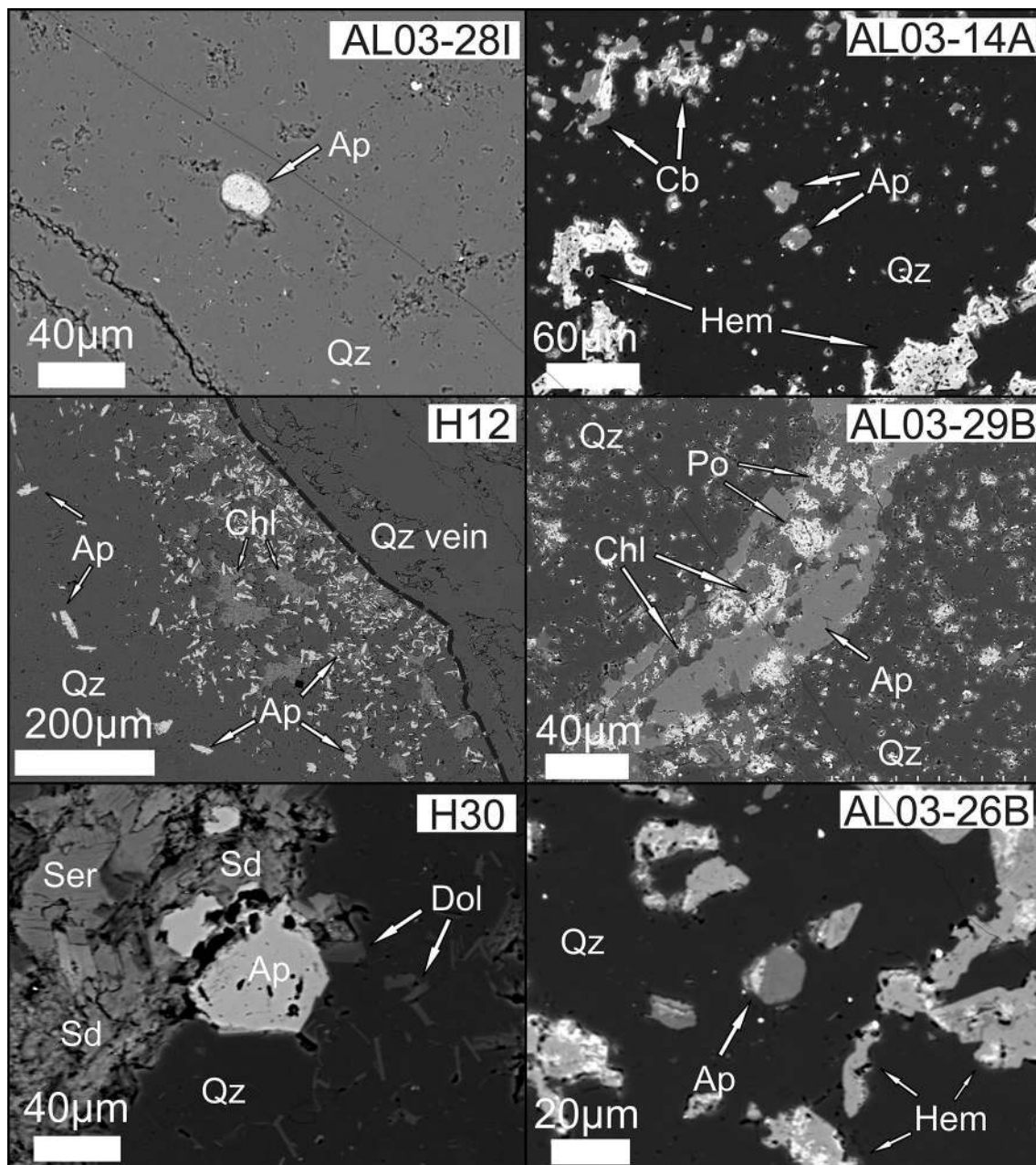


Fig. 3 Backscattered electron images of the metachert (AL03-28I, AL03-14A, and H12), igneous-derived units (H30 and AL03-29B), and BIF (AL03-26B) samples collected from the BGB

carbonates are typically represented by an Fe-rich phase (siderite) that contains minor Mg, Mn, and Ca, but dolomite, rhodochrosite, and calcite are also present. Chlorite is the secondary phase and occurs in contact with cross-cutting veins (Fig. 3). In sample H12, apatite crystals are associated with a system of veins and are relatively small ($< 50 \mu\text{m}$) (Table 1; Fig. 3).

Sample AL03-29B is silicified felsic volcanoclastics from the Mendon Formation (Blake et al. 2010) and contains mostly quartz. Chlorite is a minor compound and occurs in cross-cutting veins (Fig. 3). These veins are a clear indication of secondary hydrothermal/metamorphic overprinting in this sample. The accessory phases are represented by magnetite, pyrite, siderite, dolomite, rhodochrosite, calcite, titanite, and zircon. The apatite crystals in sample AL03-29B are relatively large and more common (up to $100 \mu\text{m}$) and are observed in the cross-cutting chlorite veins in this sample (Table 1; Fig. 3).

Sample H30 represents dacite-derived, silicified sandstone (i.e., the erosional product of underlying dacitic volcanic rocks) from the Hooggenoeg Formation of the Onverwacht Group (Hofmann and Harris 2008). Quartz and siderite are the major phases in this sample, and hematite, dolomite, rhodochrosite, and calcite are also present (Fig. 3). In addition, some alteration minerals are observed. Sericite has developed after feldspars and is a principal Al carrier phase in sample H30. The accessory mineral assemblage includes pyrite and titanite, as well as monazite and zircon. In sample H30, the apatite crystals are randomly scattered and are up to $100 \mu\text{m}$ in size, but mainly $< 50 \mu\text{m}$ (Table 1; Fig. 3).

Sample AL03-26B is a BIF from the Fig Tree Group (Blake et al. 2010). It is composed of intercalated Fe-oxide and quartz-rich layers. Hematite, together with some magnetite, is the most common Fe-oxide, with minor goethite due to recent weathering. The minor phases are carbonates (calcite, dolomite and rhodochrosite) and pyrite. Apatite is rare in the analyzed sample but is of a size up to $50 \mu\text{m}$ and was observed in association with Fe-rich layers (Table 1; Fig. 3).

Analytical approach

Sample preparation

The apatite crystals for the Fourier transform infrared (FTIR) spectroscopy were extracted from the rock samples by multistage mineral separation: the rock fragments were crushed and sieved to obtain a fraction less than $315 \mu\text{m}$, and the finest fraction was removed by careful and repeated washing; then, the samples underwent magnetic separation, heavy liquid (tetra-bromoethane and diiodomethane)

separation and careful hand-picking under a binocular microscope. The samples were too small to orient them crystallographically, and so randomly oriented apatite crystals were polished on two opposite sides to make the two surfaces parallel. The sample thicknesses were determined using the eyepiece reticule and stage micrometer scale of an optical microscope. Based on our calibration, this procedure has an uncertainty of $\pm 5\%$. The final thickness of the doubly polished crystals varied between 20 and $40 \mu\text{m}$. Petrographic thick sections (approx. $300 \mu\text{m}$ thick) were used for the REE measurements, whereas the EMP analyses of the major and trace elements, CL observations, and preparation using FIB-milled foils for the transmission electron microscope (TEM) analyses were performed on the same crystals in petrographic-thin sections.

Cathodoluminescence (CL)

CL measurements were performed at the Institute of Mineralogy of the TU Bergakademie Freiberg on carbon-coated, polished thin sections using a “hot cathode” CL microscope HC1-LM (cf., Neuser et al. 1995). The system was operated at a 14 kV accelerating voltage and a current of 0.2 mA (current density of approximately $10 \mu\text{A}/\text{mm}^2$). Luminescence images were captured “on-line” during CL operations using a Peltier-cooled digital video camera (OLYMPUS DP72). CL spectra in the wavelength range of 380 – 1000 nm were recorded with an Acton Research SP-2356 digital triple-grating spectrograph with a Princeton Spec-10 CCD detector that was attached to the CL microscope by a silica–glass fiber guide. CL spectra were measured under standardized conditions (wavelength calibration by a Hg-halogen lamp, spot width of $30 \mu\text{m}$, and measuring time of 1 s).

Transmission electron microscope (TEM) observations

TEM analyses were conducted to determine the phosphate structure and possible presence of REE-phosphate, graphite, sulfide and/or other inclusions. Sample preparation was accomplished using focused ion beam (FIB) milling. The FIB–TEM analyses were performed at the German Research Centre for Geosciences (GFZ) in Potsdam, Germany. Areas of $20 \times 20 \mu\text{m}$ were selected in each sample for analysis. First, the samples were coated with a $1.5 \mu\text{m}$ protective layer of Pt to prevent sample damage caused by FIB sputtering and Ga-ion implantation. Then, material was sputtered out from the frontal and lateral sides of the region with the FIB operating at an accelerating voltage of 30 kV and a beam current of 30 nA . The frontal surface was then “polished” with a 3-nA FIB beam current. The final size of the TEM-foils cut directly from the grains in the thin sections was approximately $20 \times 10 \times 0.15 \mu\text{m}$. The TEM-ready foils were

placed on a perforated carbon film on a copper grid. No carbon coating was required to prevent charging under the electron beam. A detailed description of the sample preparation was published by Wirth (2004, 2009). The TEM observations were performed using an FEI Tecnai G2 F20 X-Twin TEM with a Schottky field emitter as an electron source. The TEM was equipped with a Fishione high-angle annular dark field detector (HAADF), an EDAX X-ray analyzer and a Gatan electron energy-loss spectrometer (EELS). High-resolution lattice fringe images were used to calculate the diffraction pattern (fast Fourier transform, FFT) of the host apatite as well as solid inclusions within its structure and subsequently identify them. The observed d-spacings and angles between adjacent planes were compared with the d-spacings and angles calculated from literature data. The error in the observed d-spacings is within the third decimal place of the measurement. The angles between adjacent planes must match within 1° for positive phase identification. The error in the angular measurements from the FFT was $<0.5^\circ$.

Electron microprobe analysis (EMPA)

The quantitative chemical analysis of the apatite grains was performed using a JEOL SuperProbe JXA-8230 microprobe at the Laboratory of Critical Elements AGH-KGHM (Cracow, Poland). The samples were coated with a 20 nm thick carbon film. To minimize the known potential problems associated with the EMP analysis of F (predominantly) and Cl in apatite, i.e., crystallographic orientation relationship and apatite F and Cl X-ray intensities driven by the diffusion of halogen ions due to primary electron beam operation (Stormer et al. 1993; Pyle et al. 2002), the samples were analyzed using two protocols. The first protocol, proposed by Słaby et al. (2016) and Wudarska et al. (2018), consisted of an acceleration voltage of 15 kV, a beam current of 5 nA and a spot size of 8 μm . The following natural and synthetic standards, corresponding spectral lines, peak counting times and used crystals were applied: fluorite (F K α , 10 s, LDE1), albite (Na K α , 20 s, TAPH), fluorapatite (Ca K α , 10 s, PETJ and P K α , 20 s, PETL), YPO₄ (Y L α , 10 s, PETJ), CePO₄ (Ce L α , 20 s, LIFH), tugtupite (Cl K α , 10 s, PETH), fayalite (Fe K α , 10 s, LIFL) and kyanite (Al K α , 20 s, TAPH). Tests have shown that the detection limits obtained with this protocol are relatively high, and we were therefore unable to track the concentrations of the trace elements in the apatite structure and to estimate the mean formula of the apatite. Additionally, the earlier-mentioned halogen diffusion was undetectable using this protocol. The second protocol consisted of an acceleration voltage of 15 kV, a beam current of 20 nA and a spot size of 5 μm . The natural and synthetic standards, corresponding spectral lines, peak counting times and used crystals were as follows: fluorite (F K α , 10 s,

LDE1), albite (Si K α , 20 s, TAP and Na K α , 20 s), SrSO₄ (Sr L α , 20 s, TAP), fluorapatite (Ca K α , 10 s, PETH and P K α , 20 s, PETH), anhydrite (S K α , 20 s, PETJ), YPO₄ (Y L α , 10 s, PETJ), CePO₄ (Ce L α , 20 s, LIFH), barite (Ba L α , 10 s, PETJ), tugtupite (Cl K α , 10 s, PETL), fayalite (Fe K α , 10 s, LIFL), rhodonite (Mn K α , 10 s, LIFL), kyanite (Al K α , 20 s, TAP), diopside (Mg K α , 20 s, TAP), LaPO₄ (La L α , 20 s, LIF), NdPO₄ (Nd L α , 20 s, LIFL) and GdPO₄ (Gd L α , 20 s, LIFH). The apatite formulae were calculated based on the EMPA data using the method by Ketcham (2015) and were normalized based on 25 oxygen equivalents. The concentrations of all components were compared to the detection limits (DL) of each element, and only those results above the DL were taken into consideration. The OH content was calculated with the assumption that the X-site is fully occupied by F, Cl and OH (F + Cl + OH = 2).

Laser ablation inductively coupled plasma mass spectrometry (LA-ICP-MS)

For trace element analyses, LA-ICP-MS was carried out at the Geochemisches Institute, University of Göttingen, using an Element 2 (Thermo-Fischer) instrument combined with a RESolution Laser (193 nm ArF Excimer) by Resonetics equipped with the Laurin S-155 sample cell. Single measurements with a spot size of 23 μm were conducted on the apatite crystals in low mass resolution mode, and ablation was performed in a helium atmosphere. The total measurement time was typically 100 s: the signal was collected for 35 s from the target mineral, and the measurements started after 40 s of prewarming and ended after 20 s of delay. This protocol was intended to avoid peaks that do not reveal the actual concentrations. The laser was operated at a pulse energy of 80 mJ, a fluence of 2–3 J/cm² and a repetition rate of 7 Hz. The measured intensities were transformed into concentrations by applying an external calibration with NIST 610 and 612 glass reference materials. The calcium concentration was used as an internal standard. The mass spectra were visualized with Iolite v2.1528, a noncommercial freeware solution implemented as a self-contained package running in the Igor Pro v6.34A (Wavemetrics Incorporated, Portland, Oregon, USA) environment. The concentration values were obtained by integrating the results over defined time intervals. The rare earth element (REE) distribution profiles were normalized to chondritic abundances using values given by Evensen et al. (1978).

Fourier transform infrared (FTIR) spectroscopy

Unpolarized FTIR spectra across the OH-stretching range were collected using a Bruker Hyperion 2000 IR-microscope attached to a Vertex 80v FTIR-spectrometer at the German Research Centre for Geosciences (GFZ) in

Potsdam, Germany. We used a CaF₂ beamsplitter and an InSb detector and placed the crystals on KBr pellets. The reference spectra were taken next to the crystals through the KBr. The spectra were averaged over 512 scans with a spectral resolution of 2/cm. The peak separation and peak area A_i (/cm) were fitted using PeakFit software, and the bands were assigned according to Tacker (2004). Using unpolarized spectra, we can only estimate the water content, as the quantification of the water content by FTIR spectroscopy requires the measurement of polarized spectra on two orthogonal sections oriented parallel to each of two axes of the indicatrix ellipsoid (Libowitzky and Rossman 1996). We averaged the thickness-normalized integrated absorbance values measured on several crystals of each sample and multiplied the average by 2 to get the total integrated absorbance values. This procedure is based on the fact that the OH-vector in apatite is aligned parallel the crystallographic *c* axis, and the OH-bands yield maximum absorbance in the spectra parallel to the *c* axis and zero absorbance perpendicular. Multiplying the absorbance values of the unpolarized spectra by 3 as done for biaxial crystals (see discussion in Qiu et al. 2018) would overestimate the water content. To estimate the hydrogen content, we applied the Beer–Lambert law (with a density for apatite of 3.1 g cm³) and used the absorption coefficient as determined by Wang et al. (2011). Based on our experience with polarized and unpolarized measurements, this procedure gives the water content with an uncertainty of ± 30%.

Results

Metacherts (samples AL03-28I, AL03-14A and H12)

CL data

The apatite from the metacherts show a wide range of luminescence values. For sample AL03-28I, the apatite shows weak, patchy greenish luminescence in the core and very strong orange luminescence in the narrow rim (Fig. 4f). The luminescence is activated by manganese (Fig. 4c). The inhomogeneity is not visible under the polarizing microscope and is most likely related to secondary alterations. Sample AL03-14A shows weak luminescence (Fig. 5g), yet it is violet and activated by REE (Dy³⁺, Tb³⁺, Sm³⁺, Nd³⁺, and Eu³⁺) substitution. The collected CL spectrum shows very strong and sharp REE³⁺ emission bands (Fig. 5c). The CL images of sample H12 reveal that the apatite crystals are generally patchy, with orange-brownish luminescence (Fig. 6f) related to the presence of REEs (Dy³⁺, Tb³⁺, Sm³⁺, Nd³⁺, and Eu³⁺) and dominant Mn²⁺ emission (Fig. 6c).

TEM analysis

Three foils (no. 4332, sample AL03-28I; no. 4803, sample AL03-14A; and no. 4831, sample H12) were cut for the TEM analysis. Zones with abundant nanopores were observed in all three samples (Figs. 4d, e, 5d, 6d, e). Moreover, highly elongated iron sulfide inclusions were observed in samples AL03-28I and AL03-14A (Fig. 4d, e and 5d–f). The diffraction patterns (FFT) from the high-resolution lattice fringe image of sample AL03-28I were used to determine the structural parameters (d_{hkl}) of the inclusions and to identify the iron sulfide phase. The structural parameters indicate that the iron sulfide phase is pyrrhotite (Figs. 4e, 5d, f). Furthermore, silica, rutile and carbonate mineral inclusions have been identified (Fig. 4e).

In sample H12, the apatite is composed of slightly mis-oriented subgrains that show individual diffraction contrast patterns when tilting the foil in the TEM. The subgrains are separated by arrays of dislocations that form low-angle grain boundaries (Fig. 6d). Some crystal parts are porous (Fig. 6d), whereas others are rich in nanosize inclusions (Fig. 6e). The presence of highly elongated CaYP₇O₂₀ inclusions, with lengths of up to 300 nm and widths of approximately 15 nm, was identified. A more detailed description can be found in Birski et al. (2018).

Electron microprobe analysis

Apatite crystals were analyzed in three metachert samples (AL03-28I, AL03-14A and H12). The EMP data indicate that the apatite from metacherts can be divided into two groups with different volatile compositions. The first group (samples AL03-28I and AL03-14A; Figs. 4a, 5a) is fluorapatite with a significant but variable amount of OH, and the second group (sample H12) is fluorapatite (Fig. 6a). In general, both groups are Cl-poor varieties. The OH-rich fluorapatite contains substantial concentrations of S (0.28–1.11 wt%), Si (up to 1.02 wt%), Y and REEs (up to 1.72 wt%), whereas the fluorapatite is generally depleted in all trace elements (Supplementary Tables S1 and S2). The calculated OH content (assuming that the X-site is completely filled, i.e., F + Cl + OH = 2) is in the range of 30.4–40.6 mol% for sample AL03-28I, 13.9–28.0 mol% for sample AL03-14A and up to 5.7 mol% for sample H12. This is equal to 0.53–0.71, 0.24–0.49 and up to 0.10 wt% H₂O, respectively (Table 2).

Trace element composition

The REE patterns of the apatite from sample AL03-28I are flat, and europium shows a distinct positive anomaly (Fig. 4b). In addition, the AL03-14A REE pattern is relatively flat and characterized by a positive Eu, yet the anomaly

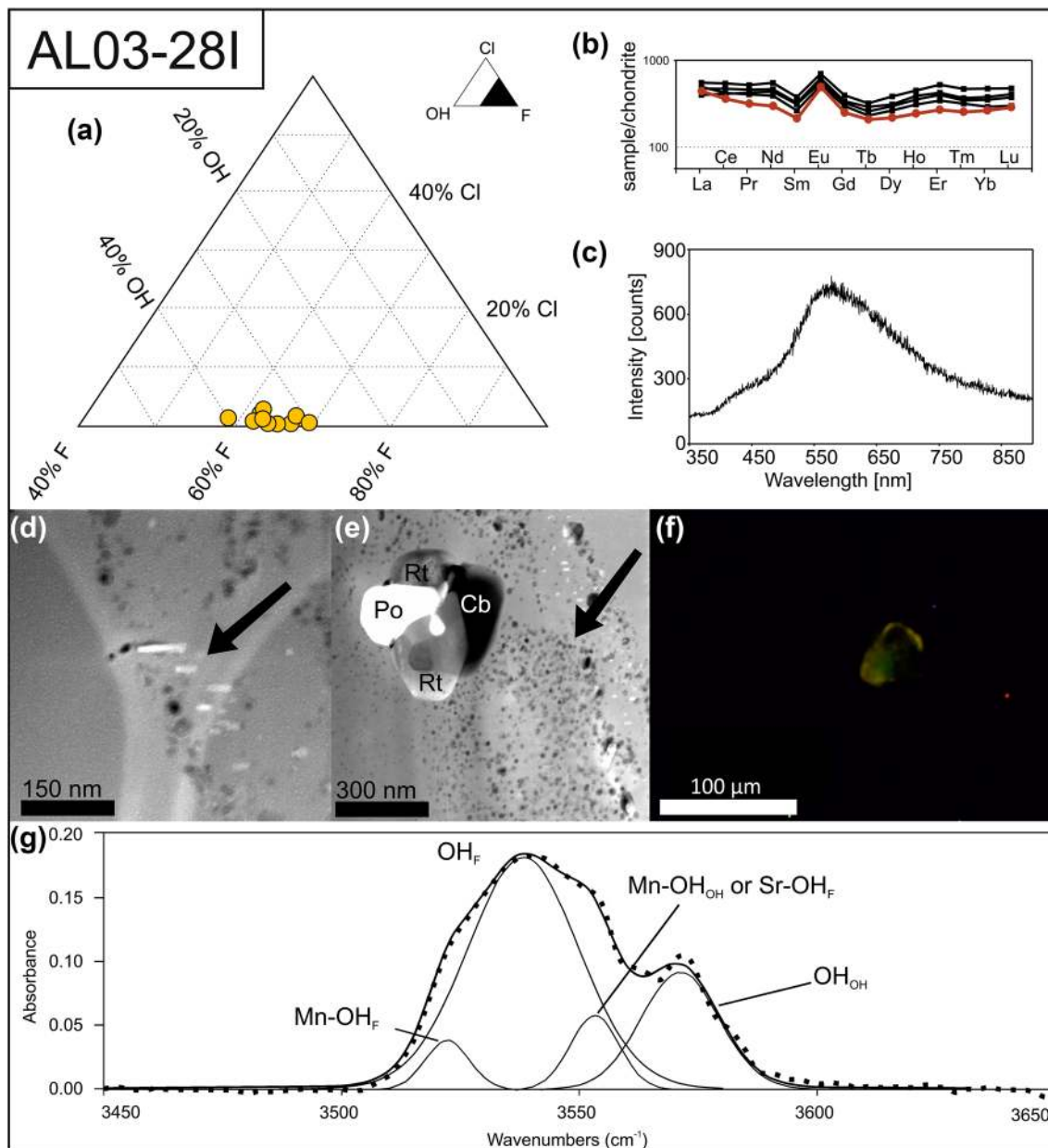


Fig. 4 Data collected for sample AL03-28I. **a** Results of electron microprobe measurements of apatite plotted in F-OH-Cl ternary diagram representing mole fraction occupancy of the X-site. **b** chondrite-normalized (Evensen et al. 1978) rare earth elements distribution patterns. Red line corresponds to REE distribution pattern of Archean sedimentary apatite (Lepland et al. 2002). **c** CL spectrum

of apatite in Fig. 4f. **d** HAADF image of apatite. Highly elongated and preferentially oriented pyrrhotite inclusions are marked (black arrow). **e** HAADF image of a rutile (Rt), carbonate minerals (Cb) and pyrrhotite (Po) inclusions in apatite. Furthermore, nano-size pores are visible (black arrow). **f** CL image of apatite. **g** FTIR spectrum

is not as distinctive as that in sample AL03-28I (Fig. 5b). Furthermore, in contrast to AL03-28I, some crystals show low LREE values. The REE patterns of both samples do not differ from those considered as sedimentary (Lepland et al. 2002; Wudarska et al. 2018).

Based on the abundance of REEs, two populations (A and B) of apatite can be distinguished in H12 (Fig. 6b). Both populations of apatite are LREE-depleted and are

characterized by similar LREE concentrations. They show increasing amounts of MREEs without any distinct anomaly. One of the patterns from population B displays a weak, positive Sm anomaly. Furthermore, the apatite grains from population B are more HREE-depleted than the apatite grains from population A. Depletion in both LREEs and HREEs is typical for metasomatic apatite (e.g., Lepland et al. 2002; Wudarska et al. 2018). The Y-Ho ratios for samples

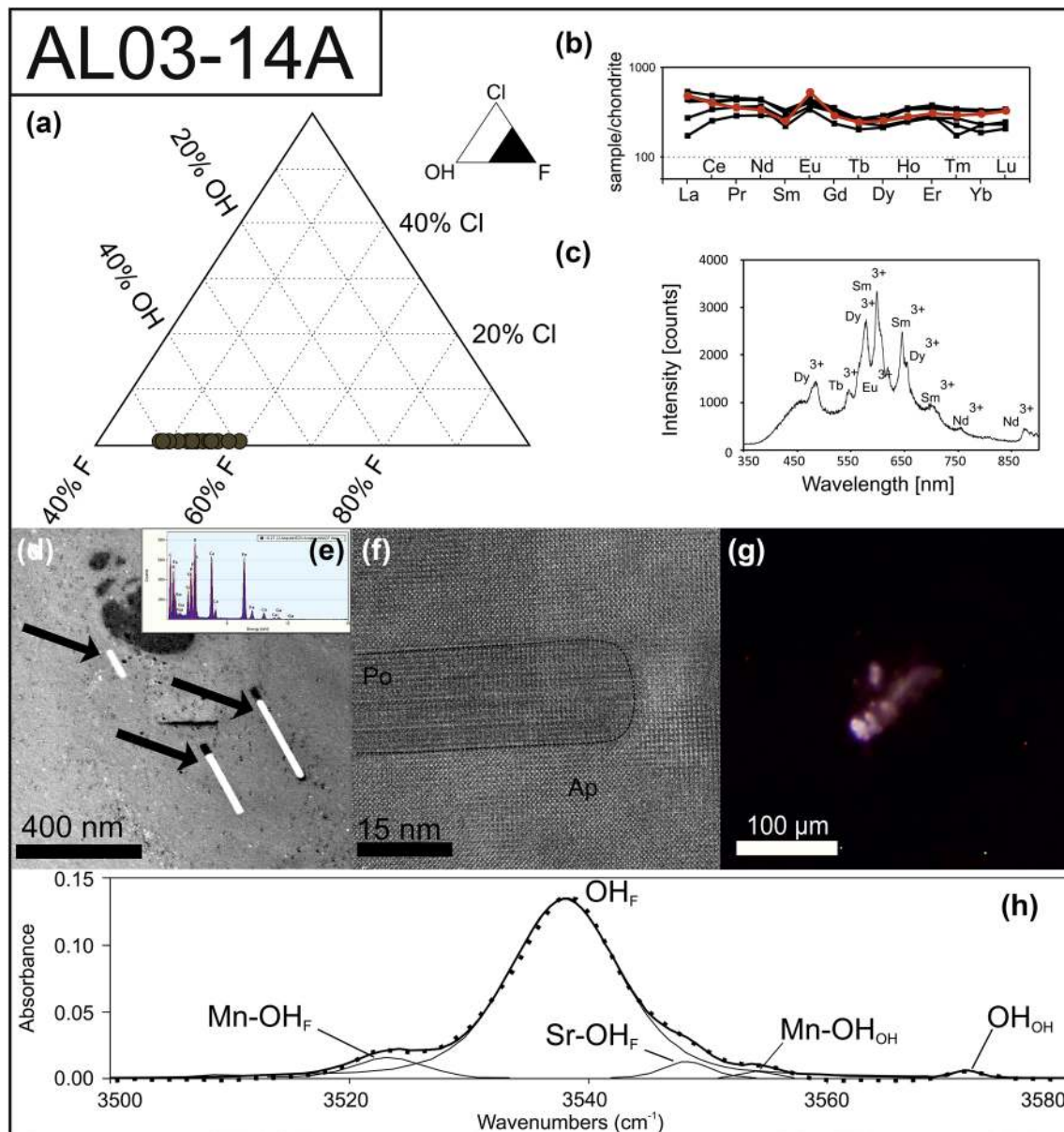


Fig. 5 Data collected for sample AL03-14A. **a** Results of electron microprobe measurements of apatite plotted in F–OH–Cl ternary diagram representing mole fraction occupancy of the X-site. **b** chondrite-normalized (Evensen et al. 1978) rare earth elements distribution patterns. Red line corresponds to REE distribution pattern of Archean sedimentary apatite (Lepland et al. 2002). **c** CL spectrum

of apatite in **g**. **d** HAADF image of highly elongated and preferentially oriented pyrrhotite inclusions (black arrow) in host apatite. **e** EDX analysis of pyrrhotite inclusions. Interference from host apatite is visible as Ca and P peaks. **f** HREM image of pyrrhotite (Po) inclusions in host apatite (Ap). **g** CL image of apatite in sample AL03-14A. **h** FTIR spectrum

AL03-28I and AL03-14A have ranges of 54–58 and 64–66, respectively, whereas sample H12 has a low Y–Ho ratio in the range of 25–52. No differences in the Y–Ho ratio have been observed between the A and B populations. The ratio values for samples AL03-28I and AL03-14A are characteristic of modern seawater (Bau and Dulski 1994), whereas the values for H12 have a broad range between the chondrite and modern seawater values (Bau and Möller 1992; Bau and

Dulski 1994). A full data set for samples AL03-28I, AL03-14A and H12 is included in Supplementary Table S3.

FTIR spectroscopy

The FTIR spectra collected from the metachert samples (Figs. 4g, 5g, 6g) show prominent OH_F bands (Tacker 2004) at 3538/cm, 3538/cm and 3545/cm in samples AL03-28I, AL03-14A and H12, respectively (Table 3). Additionally,

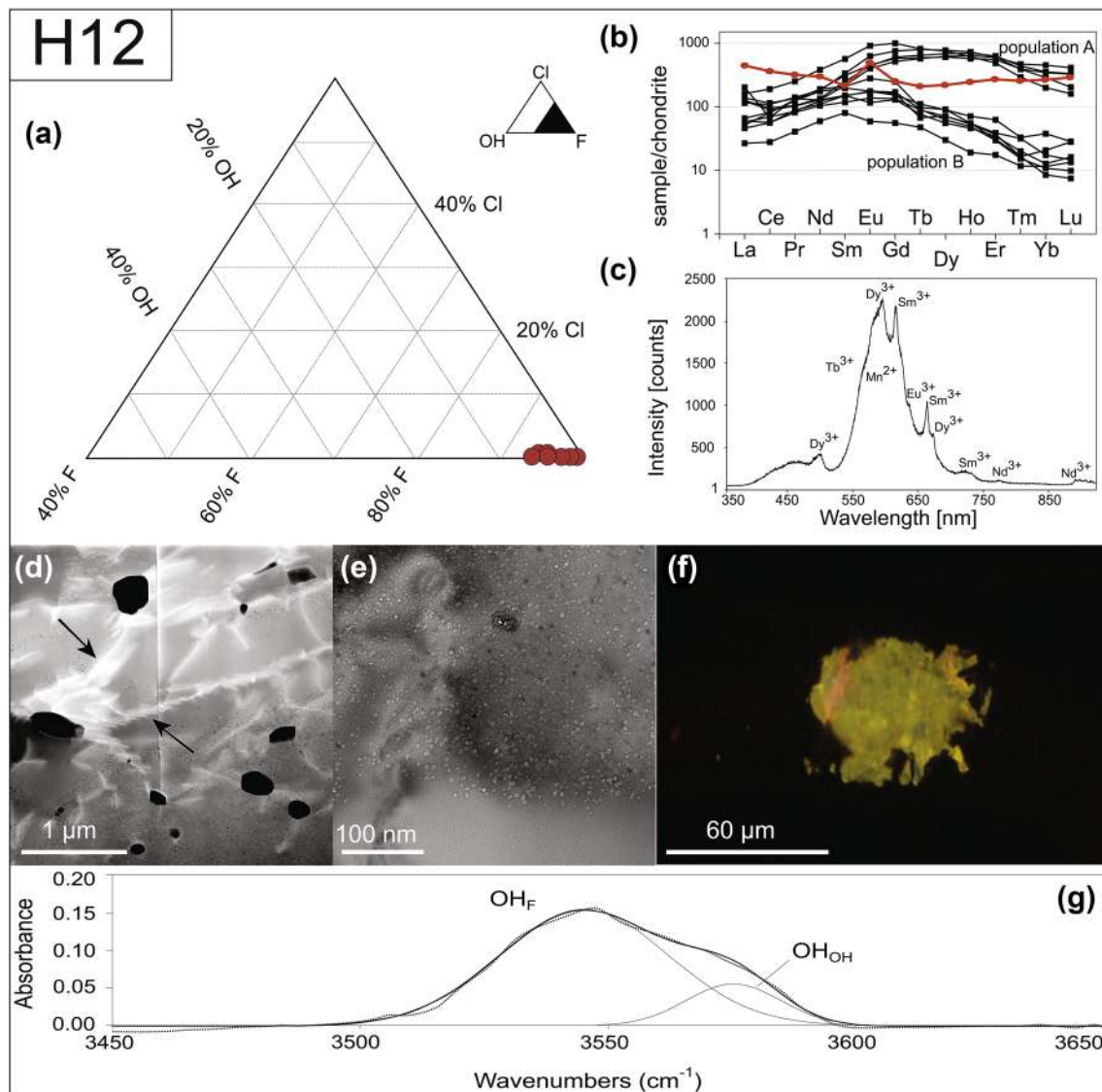


Fig. 6 Data collected for sample H12. **a** Results of electron microprobe measurements of apatite plotted in F–OH–Cl ternary diagram representing mole fraction occupancy of the X-site. **b** chondrite-normalized (Evensen et al. 1978) rare earth elements distribution patterns. Red line corresponds to REE distribution pattern of Archaean

sedimentary apatite (Lepland et al. 2002). **c** CL spectrum of apatite in **f**. **d** HAADF image of porosity in apatite. Low-angle grain boundaries are marked with black arrows. **e** Bright field image of nano-size inclusion rich zone. **f** CL image of apatite in sample H12. **g** FTIR spectrum

in all three samples, bands indexed as OH_{OH} peaks were observed, and their positions are 3572/cm, 3572/cm and 3576/cm, respectively (Table 3). No additional peaks were observed in sample H12 (Fig. 6g), whereas in samples AL03-28I and AL03-14A, additional weak bands were marked, namely, four bands (Mn-OH_{F} , Sr-OH_{F} , Mn-OH_{OH} and Sr-OH_{OH}) in sample AL03-28I (Fig. 4g) and two bands (Mn-OH_{F} and an additional band indexable as Mn-OH_{OH} or Sr-OH_{F}) in sample AL03-14A (Fig. 5g). These peaks correlate with small amounts of Mn and Sr, which have been confirmed by EMP analyses (Supplementary Table S2). With the $A_{\text{int,tot}}$ values, we estimated the wt% H_2O in the apatite

using the calibration given by Wang et al. (2011). The values of A_{int} of 11.62/cm, 11.03/cm and 5.71/cm and thicknesses of 15.6 μm , 25.3 μm and 47.4 μm correspond to 0.56 wt% H_2O , 0.33 wt% H_2O and 0.09 wt% H_2O in samples AL03-28I, AL03-14A and H12, respectively (Table 3).

Igneous-derived units (H30 and AL03-29B)

CL data

The apatite in sample AL03-29B shows strong and patchy orange luminescence (Fig. 7f) related to the presence of

Table 2 Cl and F concentrations in Barberton apatite (with 1 sd) determined by EPMA and calculated atoms per formula unit (apfu) assuming $F + Cl + OH = 2$

Sample ID	Rock type	n^a	Volatile concentration [wt%]		Atoms per formula unit (apfu)			H_2O concentration [wt%]
			[Cl]	[F]	Cl	F	OH	
AL03-28I	Chert	8	0.07 ± 0.06	2.44 ± 0.12	0.02	1.29	0.69	0.53–0.71
AL03-14A	Chert	10	0.04 ± 0.02	2.84 ± 0.14	0.01	1.56	0.43	0.24–0.49
H12	Chert	15	0.02 ± 0.01	3.62 ± 0.06	0.00	1.96	0.03	0.00–0.10
AL03-29B	Volcano-clastic	25	0.00 ± 0.01	3.72 ± 0.09	0.00	1.98	0.02	0.00–0.13
H30	Dacite-derived sandstone	13	0.22 ± 0.13	3.56 ± 0.10	0.06	1.89	0.05	0.00–0.11
AL03-26B	BIF	21	0.01 ± 0.01	2.21 ± 0.09	0.00	1.18	0.82	0.64–0.81

^aNumber of measurements conducted on a sample

Table 3 Results of FTIR measurements showing the detected absorption bands with associated molecules

Sample ID	AL03-28I	AL03-14A	H12	AL03-29B	H30	AL03-26B
$A_{int, tot}$ [/cm]	11.62	11.03	5.71	5.91	5.29	12.84
Thickness [μm]	15.6	25.3	47.4	54.2	52.6	22.7
C_{H_2O} [wt %]	0.56	0.33	0.09	0.08	0.08	0.43
OH_F						
FWHM ^b	27.705	10.917	41.505	28.711	16.154	40.235
Position ^c	3538.6	3538.1	3545.2	3539.5	3538.4	3544.7
A_{int} [/cm]	7.20	8.79	5.45	8.75	4.47	5.19
OH_{OH}						
FWHM ^b	19.473	2.538	22.042	14.124	6.615	21.865
Position ^c	3571.6	3571.7	3575.7	3571.6	3571.2	3573.2
A_{int} [/cm]	2.46	1.02	0.25	1.84	0.14	0.72
$Mn-OH_F$						
FWHM ^b	11.102	6.912		11.679	6.563	
Position ^c	3522.5	3523.4		3522.7	3522.4	
A_{int} [/cm]	0.97	0.33		0.74	0.19	
$Mn-OH_{OH}$						
FWHM ^b	12.187 ^a	3.262		6.088	9.752 ^a	
Position ^c	3553.6 ^a	3554.2		3562.1	3554.0 ^a	
A_{int} [/cm]	1.00 ^a	0.12		0.36	0.49 ^a	
$Sr-OH_F$						
FWHM ^b		4.636		9.013		
Position ^c		3548.3		35553.5		
A_{int} [/cm]		0.77		0.82		
$Sr-OH_{OH}$						
FWHM ^b				6.717		
Position ^c				3583.9		
A_{int} [/cm]				0.32		

^aOH-band can be indexed either as $Mn-OH_{OH}$ or $Sr-OH_F$

^bFWHM full width at half maximum

^cGiven in /cm

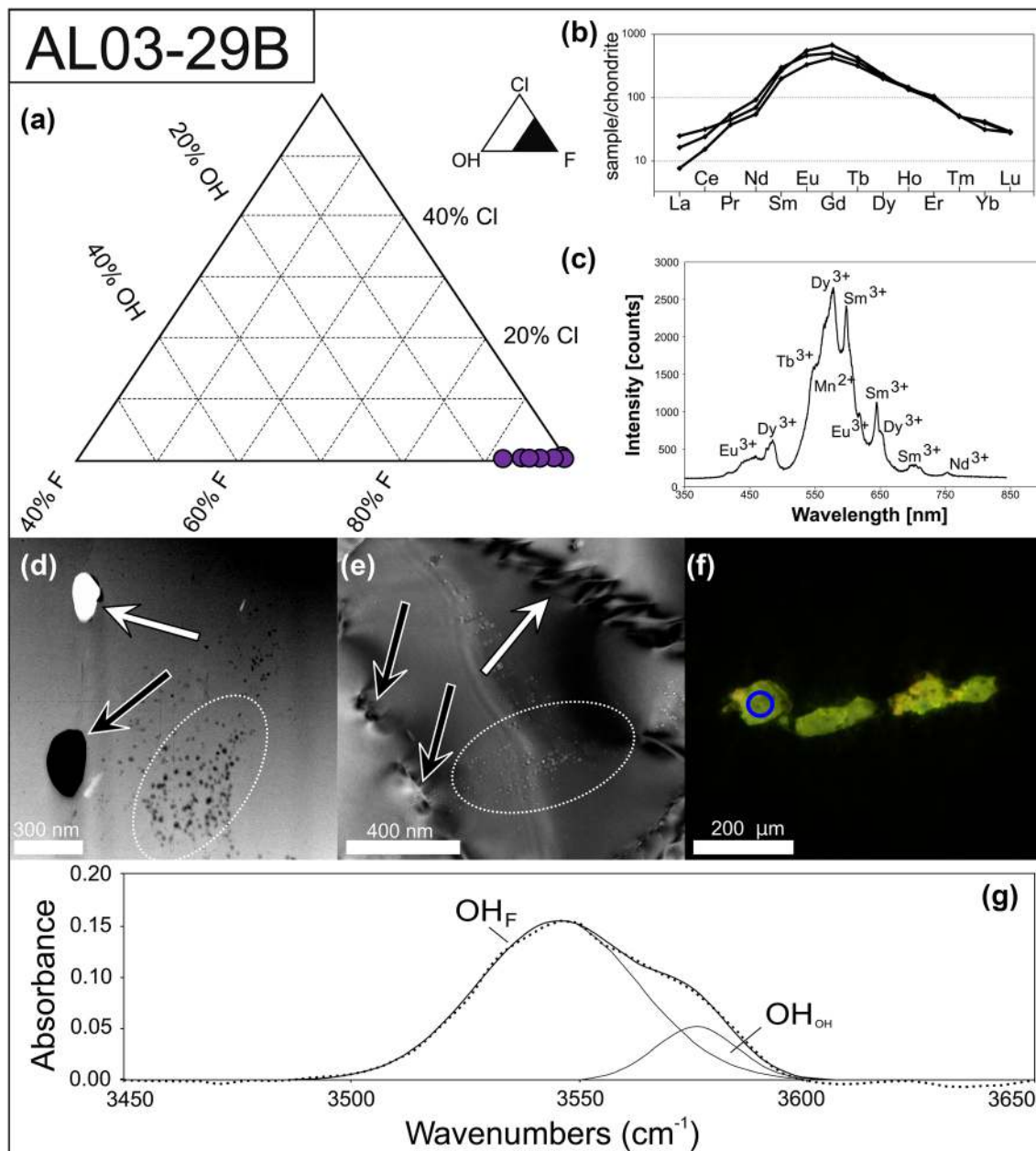


Fig. 7 Data collected for sample AL03-29B. **a** Results of electron microprobe measurements of apatite plotted in F–OH–Cl ternary diagram representing mole fraction occupancy of the X-site. **b** chondrite-normalized (Evensen et al. 1978) rare earth elements distribution patterns. **c** CL spectrum of apatite in **f**. **d** HAADF image of metasomatically altered apatite in sample AL03-29B. Nano-

size inclusion rich zone (white dots), porosity (black arrow) and Ca₃Y(PO₄)₃ inclusion (white arrow; Birski et al. 2018) are visible. **e** Bright field image of nano-size inclusion rich zone (white dots), low-angle grain boundaries (white arrow) and single dislocations (black arrows). **f** CL image of apatite in sample AL03-29B. The spectrum was collected from area marked by blue circle. **g** FTIR spectrum

Mn²⁺ and REE³⁺ (Dy³⁺, Tb³⁺, Sm³⁺, Nd³⁺, Eu³⁺) emission centers that are confirmed in the CL spectra (Fig. 7c). In sample H30, the luminescence is weak, with blue-violet luminescence in the core and very strong orange luminescence in the narrow rim (Fig. 8c). This inhomogeneity is related to secondary alterations or it could be a new growth

related to metamorphic events. The CL spectra indicate the presence of REE³⁺ and Mn²⁺ emission centers in variable amounts. The manganese emission center is dominant over bands characteristic of REE emission centers in areas that are closer to the rims (Fig. 8f).

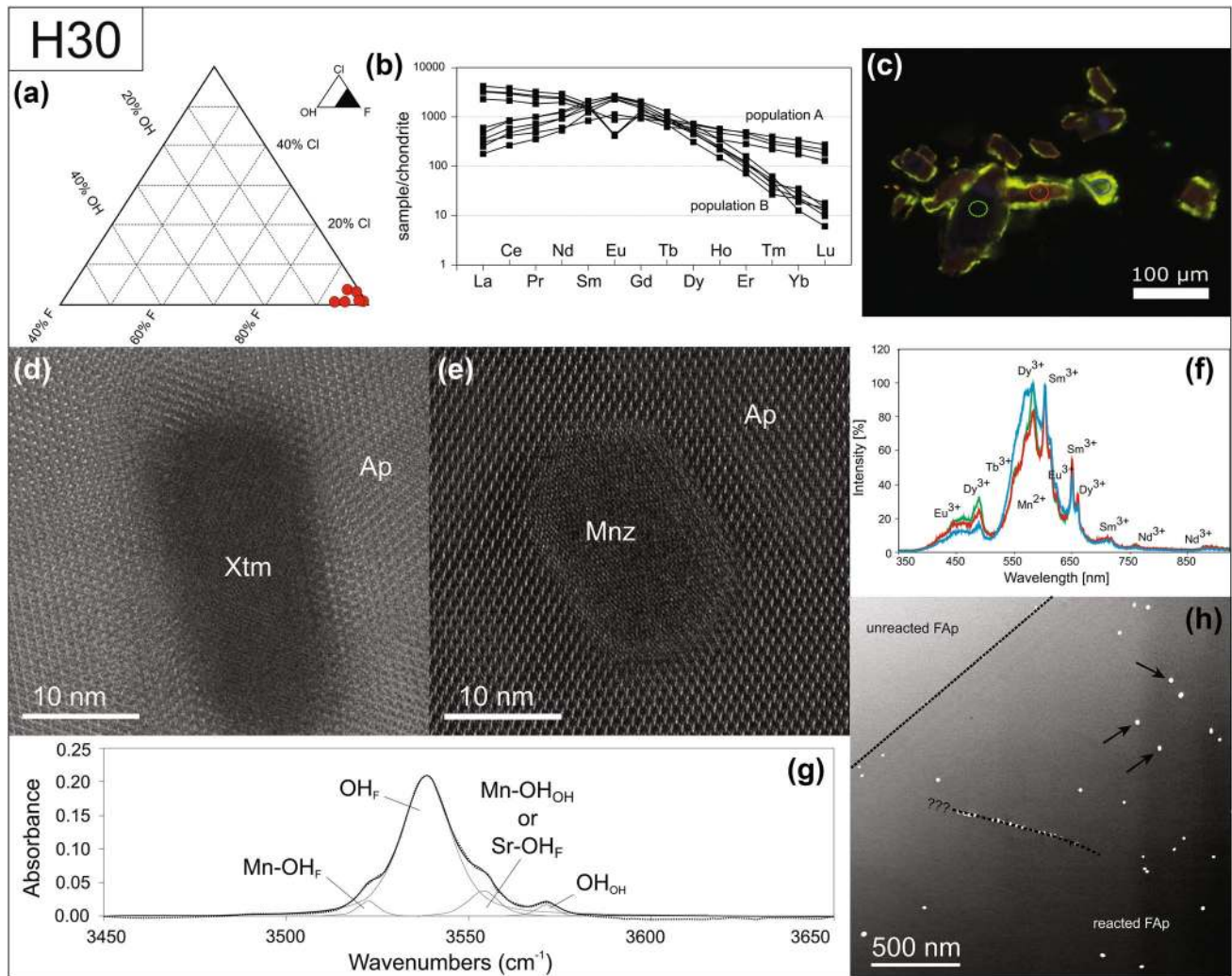


Fig. 8 Data collected for sample H30. **a** Results of electron microprobe measurements of apatite plotted in F–OH–Cl ternary diagram representing mole fraction occupancy of the X-site. **b** chondrite-normalized (Evensen et al. 1978) rare earth elements distribution patterns. **c** CL image of apatite in sample H30. **d** HREM image of xenotime inclusion in host apatite. **e** HREM image of monazite inclu-

sion in host apatite. **f** CL spectrum of apatite in **c**. Color of spectrum agrees with the area marked in **c**. **g** FTIR spectrum. **h** HAADF image of the area being the border between reacted and unreacted fluorapatite (black dashed lines). REE-phosphate inclusions are marked by black arrows

TEM analysis

Four foils, two (nos. 4332 and 4334) from AL03-29B and two (nos. 4441 and 4482) from H30, were cut. In sample AL03-29B, the apatite is composed of slightly misoriented subgrains that show individual diffraction contrast patterns (Fig. 7e). The micrometer-sized pores are partially filled with Ca₃Y(PO₄)₃ inclusions up to 200 nm in size (Fig. 7d). Furthermore, cracks, pores and nano channel-like structures have been reported by Birski et al. (2018).

Both foils from sample H30 reveal the absence of dislocations or low-angle grain boundaries in apatite, but (REE + Y)-phosphate inclusions were observed. The diffraction patterns (FFT) from the high-resolution lattice fringe

image provide information about the lengths of the individual diffraction vectors and the angles between them, allowing mineral identification. Therefore, FFTs were collected from the (REE + Y)-phosphate inclusions in both foils. In foil no. 4332, the angles between the vectors were measured in three inclusions. The first inclusion featured observed angles of 37.3° and 50.6° (Supplementary Table S4 a), the second featured observed angles of 34.0° and 66.7° (Supplementary Table S4 b), and the third featured observed angles of 28.0°, 21.0°, 81.3° and 48.7° (Supplementary Table S4 c). All three inclusions were indexed as xenotime (Supplementary Table S4, Fig. 8d). In foil no. 4334, the angles between the vectors were measured in three inclusions as well. All three inclusions featured observed angles

of approximately 40.0° and 53.0° (Supplementary Table S4) and were recognized as monazite (Supplementary Table S4, Fig. 8e). Furthermore, an orientation relationship between the monazite inclusion and host apatite was observed (Supplementary Table S4). The (010) direction of monazite is parallel to the (0001) direction of the apatite for all indexed monazite inclusions ($[010]_{\text{Mnz}} \parallel [0001]_{\text{Ap}}$). The presence of hexagonal symmetry in the monazite inclusions is also a characteristic feature of the interfaces observed in high-resolution images of sample H30 (Fig. 8e). In foil no. 4482, two zones could be discriminated (Fig. 8h). The first zone has no inclusions and is visibly brighter than the second zone, which commonly contains solid inclusions. The differences in brightness in the high-angle annular dark-field (HAADF) Z-contrast images are most likely due to changes in the relative concentration of REEs of apatite caused by the removal of REEs and Y from the host apatite structure and the later nucleation of (REE + Y)-phosphate inclusions within the visibly darker zone. The boundary between the two zones is very clear and sharp (Fig. 8h).

Electron microprobe analysis

The apatite grains from the igneous-derived units are predominantly fluorapatite (Figs. 7a, 8a). The Cl and trace element concentrations are below the detection limit (Supplementary Tables S1 and S2). The calculated OH content (assuming that the X-site is completely filled, i.e., $F + Cl + OH = 2$) is up to 7.54 mol%, which equals up to 0.13 wt% H₂O (Table 2).

Trace element composition

The REE patterns of sample H30 indicate the presence of two populations of apatite crystals (Fig. 8b). Population A is slightly enriched in LREEs and slightly depleted in HREEs and has a distinct negative Eu anomaly. This REE pattern is characteristic of magmatic apatite crystallization after plagioclase fractionation. Population B is characterized by an enrichment in MREEs and a strong depletion in LREEs and HREEs compared to population A. Population B does not demonstrate any distinct anomalies. The apatite crystals from sample AL03-29B are characterized by an enrichment in MREEs (Fig. 7b). The REE distribution patterns of population B in samples H30 and AL03-29B are typical of metasomatic apatite crystals (e.g., Lepland et al. 2002; Wudarska et al. 2018), and the Y-Ho ratio ranges between 21 and 28. A full data set is included in Supplementary Table S3.

FTIR spectroscopy

The water content was also determined in apatite from the igneous-derived samples using FTIR spectrometry.

The spectra of sample AL03-29B show the presence of two bands, a prominent OH_F band at 3539/cm and a much weaker OH_{OH} band at 3545/cm, whereas in the spectra of sample H30, we observed four OH-bands, i.e., a dominant OH_F band at 3538/cm and three additional small bands (OH_{OH}, Mn–OH_F and a band indexable as Mn–OH_{OH} or Sr–OH_F) (Table 3; Figs. 7g, 8g). These spectra have integrated absorbance values of 5.29/cm and 5.91/cm for H30 and AL03-29B, respectively. By applying the equation by Wang et al. (2011) and using measured thicknesses of 52.6 μm (H30) and 54.2 μm (AL03-29B), a hydroxyl content equivalent to approximately 0.08 wt% H₂O can be calculated for the apatite in both samples (Table 3).

Banded iron formation (sample AL03-26B)

CL data

The apatite shows very weak yellow luminescence (Fig. 9f). The collected CL spectrum indicates that this luminescence is related to Mn²⁺ substitution (Fig. 9c). The inhomogeneity is easily visible. The core is dull and almost completely dark, which is most likely caused by Fe substitution (or a limited Mn content), whereas the very narrow rim is much brighter. This heterogeneity is not visible under the polarizing microscope.

TEM analysis

One foil (no. 4829) was lifted out from the sample for TEM study. The studied apatite was relatively rich in micro- and nanosized iron oxide inclusions and pores (with sizes up to 200 nm) (Fig. 9e). Furthermore, deformation and annealing features such as parallel dislocations and low-angle grain boundaries were observed (Fig. 9d).

Electron microprobe analysis

In sample AL03-26B, the apatite is associated with iron oxide layers and it is fluorapatite (Fig. 9a), with Cl concentrations below the detection limit (Supplementary Table S1). Furthermore, the sample contains appreciable concentrations of Sr (approx. 0.27 wt%) (Supplementary Table S2). The OH content, calculated assuming that the X-site is completely filled, i.e., $F + Cl + OH = 2$, is in the range of 36.7–45.8 mol%, which equals 0.64–0.81 wt% H₂O (Table 2).

Trace element composition

The apatite crystals from sample AL03-26B are relatively enriched in REEs (Fig. 9b). Moreover, they are characterized by flat REE distribution patterns with slightly positive Nd,

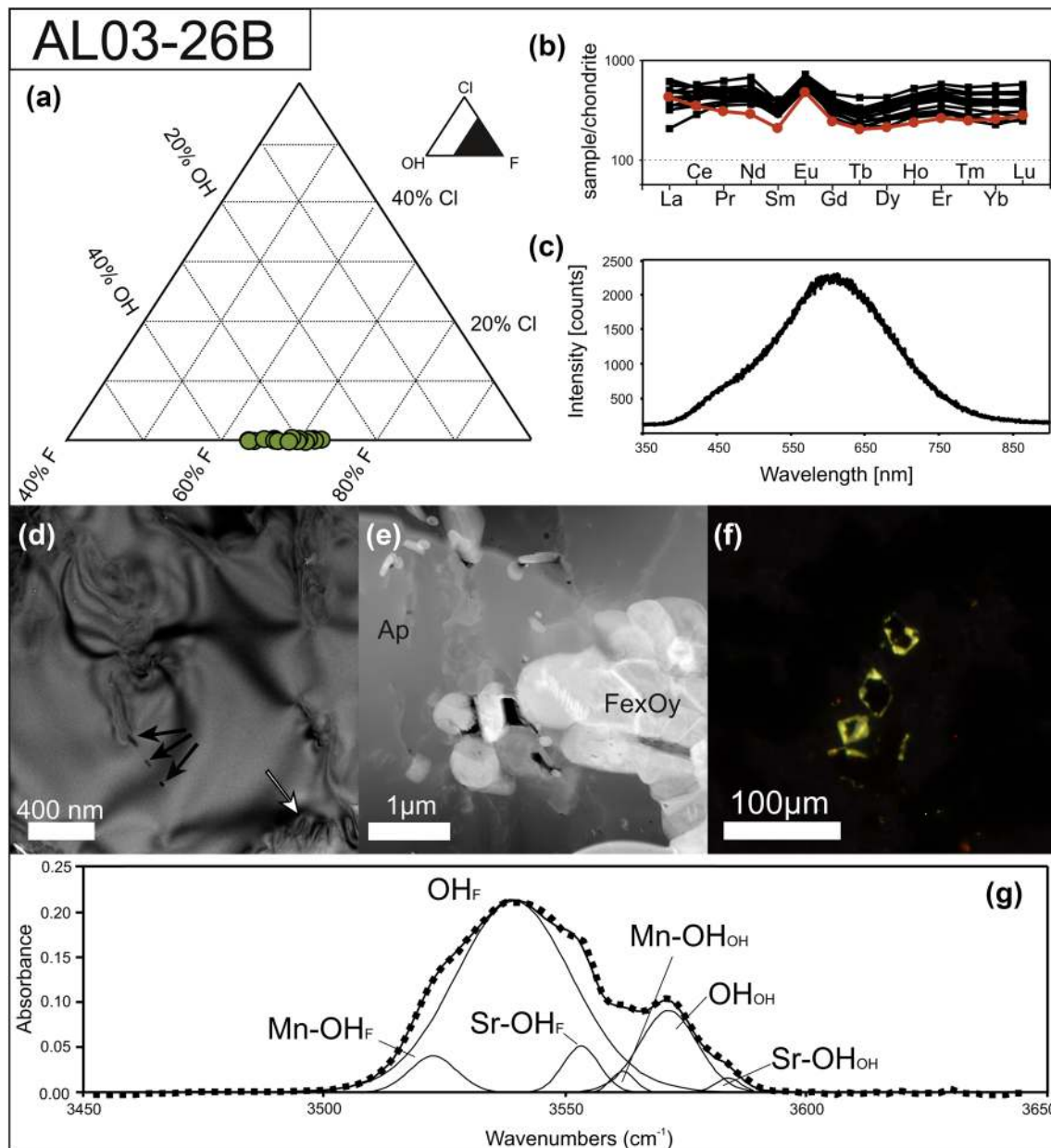


Fig. 9 Data collected for sample AL03-26B. **a** Results of electron microprobe measurements of apatite plotted in F–OH–Cl ternary diagram representing mole fraction occupancy of the X-site. **b** chondrite-normalized (Evensen et al. 1978) rare earth elements distribution patterns. Red line corresponds to REE distribution pattern of Archaean sedimentary apatite (Lepland et al. 2002). **c** CL image

of apatite in sample AL03-26B. **d** Bright field image of apatite. Iron oxide inclusions are marked with black arrows, whereas low-angle grain boundaries with white arrow. **e** HAADF image of the iron oxide inclusions in host apatite. **f** CL spectrum of apatite in Fig. 9c showing a broad emission band due to Mn^{2+} . **g** FTIR spectrum

negative Sm, and distinct positive Eu anomalies. The Y–Ho ratio varies in the range of 55–70. This REE pattern is characteristic of sedimentary apatite (e.g., Lepland et al. 2002; Wudarska et al. 2018), whereas the Y–Ho ratio is characteristic of modern seawater values (Bau and Dulski 1994). A full data set is included in Supplementary Table S3.

FTIR spectroscopy

The FTIR spectrum collected from the sample shows a multiband pattern (Fig. 9g). A prominent OH band occurs at 3539/cm (Table 3). According to Tacker (2004), the OH band corresponds to a high F content within the apatite and should be indexed as an OH_F band. Additional peaks were

observed at 3572/cm (Table 3) and should be correlated with the position of the OH_{OH} band. Four additional weak bands can be marked as Mn-OH_{F} , Sr-OH_{F} , Mn-OH_{OH} and Sr-OH_{OH} (Fig. 9g). Their presence may be correlated with the presence of small amounts of Mn and Sr, as confirmed by EMPA (Supplementary Table S2). The integrated absorbance, $A_{\text{int,tot}}$, was determined to be 12.84/cm. By applying the calibration of Wang et al. (2011) and using a measured thickness of 22.7 μm and a density of 3.1 g cm^3 , a hydroxyl content equivalent to 0.43 wt% H_2O was calculated for this sample (Table 3).

Discussion

Apatite formation/transformation pathways

The studied samples show different paths of formation and also dissimilar transformation paths. The question arises as to whether the obtained characteristics allow the confirmation of the expected environment of emergence if all samples are transformed. What part of the characteristics can be considered as an indicator for the environment of the initial formation of phosphates? Finally, given that the studied phosphates originated from the period before the Great Oxidation Event, it would be interesting to know if the turn from a prebiotic to a biotic environment somehow influenced their transformation paths.

Modern ocean waters are characterized by a positive Y anomaly (mass ratio of Y to Ho) and a negative Ce anomaly, and their REEs show a pattern of slight enrichment from light to heavy REEs. However, modern hydrothermal vent waters are characterized by a positive Eu anomaly (Bau and Dulski 1999). The larger Y anomalies (40–80) in modern ocean waters occur in open marine settings, while the smaller Y anomalies (33–40) occur in near-shore or restricted settings (De Baar et al. 1985a, b; Bau and Dulski 1994; Zhang and Nozaki 1996; Bau et al. 1997). The Y/Ho ratio also varies with the salinity, phosphate mineral solubility, and fractionation during chemical weathering, biological processes and redox cycling (Liu and Byrne 1997; Hill et al. 2000). Our understanding of the Y anomalies is somewhat limited, as many earlier studies did not include Y measurements alongside REEs (Tostevin et al. 2016).

Measurements of the REEs and Y in 2.5–3.7 Ga-old BIFs and metacherts (e.g., Bolhar et al. 2004, 2005; Nishizawa et al. 2005; Hofmann et al. 2013) as well as in apatite (Lepland et al. 2002; Nishizawa et al. 2004; Joosu et al. 2015, 2016; Wudarska et al. 2018) show that they contain “sea water fingerprints” with a Y anomaly and flat REE distribution patterns, identical to the pattern found in modern sea water (Elderfield and Greaves 1982). The Y anomaly seems to be present in geological records throughout the

entire Precambrian, both after and prior to the great oxidation event (GOE) and up to the Eoarchean (Joosu et al. 2015, 2016; Wudarska et al. 2018). However, unlike modern seawater, the sedimentary rocks prior to the GOE are characterized by a lack of a negative Ce anomaly, which implies anoxic conditions (e.g., Lepland et al. 2002; Joosu et al. 2015, 2016; Wudarska et al. 2018), whereas a positive Eu anomaly indicates significant hydrothermal input into the Archaean oceans (Veizer et al. 1989). Due to the mafic melt differentiation rules, apatite crystallizes as one of the latest phases, and therefore they exhibit a negative Eu anomaly (Puchelt and Emmermann 1976). During late diagenesis and hydrothermal recrystallization, apatite typically becomes enriched in the middle REEs (MREEs) because of their preferential substitution for Ca in the apatite crystal lattice due to their similar ionic radii (Morad and Felitsyn 2001), whereas light REEs (LREEs) and heavy REEs (HREEs) remain in the fluid. Such a crystal structure-controlled MREE enrichment mechanism is supported by the modeling studies of Reynard et al. (1999), who showed that MREE enrichment is expected during the late diagenetic recrystallization.

Apatite of sedimentary origin

The first group of apatite (AL03-28I, AL03-14A and AL03-26B) has the signature of an Archaean sedimentary environment. Phosphate oxygen data ($\delta^{18}\text{O} = 19.9\text{‰}$ and 18.1‰ in AL03-28I and AL03-26B, respectively) provided the basis for Blake et al. (2010) to infer the influence of microbial activity during phosphate precipitation. The REE distributions in the Archaean sediments have been shown to typically exhibit positive Eu anomalies (Derry and Jacobsen 1990; Lepland et al. 2002; Nishizawa et al. 2004; Wudarska et al. 2018), suggesting that hydrothermal venting was the major contributor to the REE budget of the Archaean ocean (Danielson et al. 1992; Lepland et al. 2002; Wudarska et al. 2018). These data are consistent with the higher geothermal gradient in the early Earth (Ronde et al. 1997). Since the crustal differentiation was probably less evolved during the Archaean and less continental crust was available for weathering, maintaining the hydrothermal REE signature of the ocean against the contribution from the continental source was easier. Furthermore, the Eu anomalies recorded in authigenic apatite are considered stable tracers that do not change during diagenesis (Shields and Stille 2001). Therefore, the nucleation of apatite in equilibrium with the Archaean seawater in AL03-28I, AL03-14A and AL03-26B is indicated by both the Y–Ho ratio (Fig. 10) and REE distribution patterns with a positive Eu anomaly (Figs. 4b, 5b, 9b). Furthermore, the apatite in all three samples is characterized by a relatively high water concentration, as shown by EMP analyses (Table 2) and FTIR (Table 3). The obtained EMPA values are in good agreement with FTIR measurements

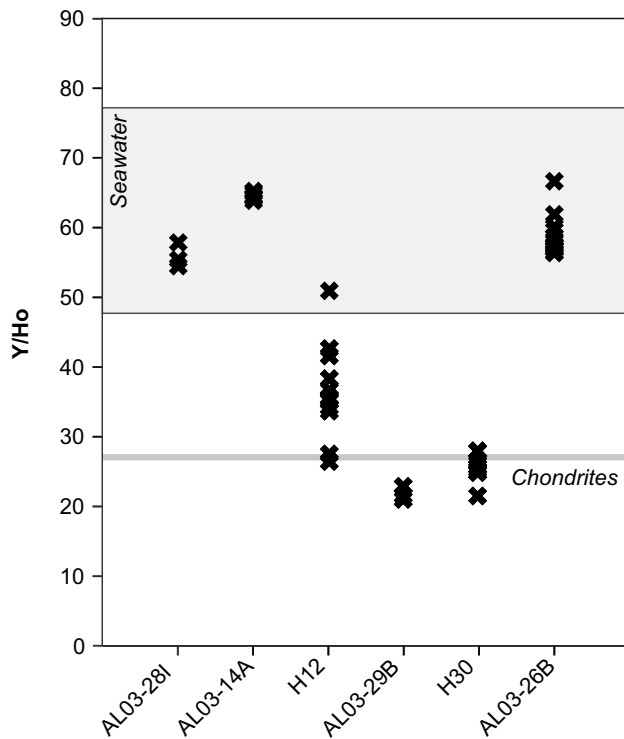


Fig. 10 Y/Ho ratios of apatite from Barberton. Ratios of the samples AL03-28I, AL03-14A and H12 are similar to the values determined for seawater (Bau and Dulski 1994). It suggests sedimentary origin of apatite. Chondritic ratio is at lower level: Y/Ho = 26–28 (Bau and Möller 1992)

(Tables 2, 3). All the values plotted in the ternary diagram indicate that the samples have a wide range of F- and OH-members (Figs. 4a, 5a, 9a). The TEM observations of these samples show no indications of secondary alteration, which allows treating a number of observed features as primary. The presence of zones rich in nanopores (Figs. 4e, 5f) is most likely associated with the precipitation process and is in agreement with the presence of highly elongated and preferentially oriented pyrrhotite inclusions (Figs. 4e–f, 5e–f). Similar features were previously observed by Gottesmann and Wirth (1997) in apatites of magmatic plutons from Kirchberg and Bergen in the Vogtland region of Saxony, Germany. The presence of epitactically intergrown apatite and pyrrhotite inclusions can be caused by their similar lattice spacing. This similarity may result in the observed orientation relationship. The nuclei of pyrrhotite formed due to the oversaturation of Fe and S ions at the interface of the seawater/phosphate precursors using the apatite lattice as a template for nucleation. Subsequently, they grew simultaneously with the apatite until the concentrations of Fe and S at the interface decreased enough to prevent the further growth of pyrrhotite. During the growth of apatite, the moving interface experienced increases in the Fe and S concentrations, thus inducing the formation of new pyrrhotite

nuclei at the interface of the seawater/phosphate precursors. The epitactic intergrowths might also explain the lack of an amorphous rim at the phase boundary interface between the pyrrhotite inclusion and the host apatite. The CL data (Fig. 5c, d) indicate that the luminescence of the apatite crystals is activated by REE substitution, which may point to the original composition of the formation environment. Archaean ocean pH value is still a matter of debate and controversy. Proposed values are scant and conflicting. Studies have concluded that the early ocean was alkaline (Kempe and Degens 1985; Blättler et al. 2017), acidic (Sleep and Zahnle 2001; Halevy and Bachan 2017; Krissansen-Totton et al. 2018), or approximately neutral (Macleod et al. 1994; Friend et al. 2008). Nevertheless, most proposed values is in the range of 6.5–8.6 pH. This lilac/violet CL color seems to be related to alkaline conditions that favor the incorporation of REEs (Kempe and Götze 2002). CL data are in the support of circumneutral to weakly alkaline Archaean ocean pH based on rare Earth element anomalies proposed by Friend et al. (2008). These assumptions correlate with determined trace element abundances (Supplementary Table S3). On the other hand, the presence of orange luminescence (activated by Mn^{2+}) in the narrow rim of sample AL03-28I (Fig. 4c) may indicate that this sample experienced secondary transformation processes and Mn substitution to a limited extent. No evidence of metamorphic growth of apatite have been observed in CL. The lack of CL luminescence (Fig. 9f) in AL03-26B is most likely caused by Fe^{2+} substitution (Kempe and Götze 2002), which is confirmed by the presence of relatively high concentrations of iron in the EMP analyses (Supplementary Table S2). Moreover, the iron concentrations in apatite correlate with the presence of iron oxide inclusions (Fig. 9e) and are associated with Fe-rich layers in the BIFs (Fig. 3). The characteristics of the samples of the first group allow them to be placed in the environment before the GOE as samples that at least partially preserve their original structures and compositions.

Metasomatic/secondary apatite

The samples from the igneous-derived rocks (AL03-29B and population B in H30) and metachert (H12) show very distinct features that are characteristic of metasomatic apatite. Blake et al. (2010) reported a $\delta^{18}O$ value of 9.3‰ for apatite crystals from the same location as sample AL03-29B. This relatively low $\delta^{18}O$ value, together with evidence for secondary veining, indicates that these apatites formed from secondary fluids (Blake et al. 2010). The result obtained in this paper provides additional evidence for this. Secondary overprinting is indicated by the metasomatic-type REE distribution pattern and Y–Ho ratio values that are much lower than the characteristic values for modern seawater (Fig. 10) (Bau and Möller 1992; Bau and Dulski 1994). Due to the

removal of LREEs and HREEs from the structure of the apatite, MREE enrichment is considered the most important characteristic to evaluate the effects of the late diagenetic alteration of apatite. This feature is clearly visible in the apatite within this group (Figs. 6b, 7b, 8b). Previous studies (Harlov et al. 2002; Harlov and Förster 2003; Wudarska et al. 2018) have shown that during metasomatic transformations, apatite evolves towards the F-rich end-member, and therefore the secondary apatite will be relatively depleted in water in comparison to the sedimentary apatite (Tables 2, 3). Furthermore, relative to the sedimentary apatite, the metasomatic apatite also seems to be depleted in trace elements due to their removal by fluids, consistent with the results for H12 from EMP analyses (Supplementary Table S2) and FTIR spectra (Fig. 6g). The features of the metasomatic influence in samples H12, H30 and AL03-29B are also very distinct under the TEM. Traces of secondary alterations have already been described (Birski et al. 2018), and they consist of slightly misoriented subgrains that are separated by arrays of dislocations forming low-angle grain boundaries, the presence of porosity, and solid inclusions that are associated with the system of subgrain boundaries (Figs. 6d, 7d, e). The presence of numerous inclusions ($\text{Ca}_3\text{Y}(\text{PO}_4)_3$ and $\text{CaYP}_7\text{O}_{20}$) in metasomatic apatite has also been reported (Birski et al. 2018), whereas in sample H30, inclusions of monazite and xenotime have been observed. The occurrence of monazite [(Ce,LREE)(PO_4)] and/or xenotime [(Y,HREE)(PO_4)] as inclusions in fluorapatite [$\text{Ca}_{10}(\text{PO}_8)_6(\text{F,Cl,OH})_2$] has been widely documented (Harlov and Förster 2003; Harlov et al. 2005; Harlov 2011, 2015; Jonsson et al. 2016). Monazite and xenotime inclusion formation has been directly linked to metasomatism by either external or internal H_2O -bearing fluids (Harlov 2015). The fluid-induced formation of monazite and xenotime inclusions in apatite is supported by experiments, which indicate that their formation and growth are the result of coupled dissolution–reprecipitation processes during metasomatic alteration (Harlov et al. 2002, 2005; Harlov and Förster 2003; Harlov 2011, 2015). In the apatite structure, (REE + Y) substitutions are charge-balanced through coupled substitutions: $\text{Si}^{4+} + (\text{REE} + \text{Y})^{3+} = \text{P}^{5+} + \text{Ca}^{2+}$ and $\text{Na}^+ + (\text{REE} + \text{Y})^{3+} = 2\text{Ca}^{2+}$ (Pan and Fleet 2002). The nucleation of the REE-phosphate inclusions in the host apatite structure is caused by fluids that preferentially remove Na and/or Si from the apatite–fluid system without removing REEs. Consequently, when supersaturated conditions are reached, the REEs can react with phosphate and possibly cause the nucleation and growth of (REE + Y)-phosphate phases. This process occurs via the two following mass transfer reactions: $[\text{Ca}_{5-2x}, \text{Na}_x, \text{REE}_x]\text{P}_3\text{O}_{12}\text{F} + x [2 \text{Ca}^{2+} + \text{P}^{5+}]$ (from the fluid) = $\text{Ca}_5\text{P}_3\text{O}_{12}\text{F} + x \text{REEPO}_4 + x[\text{Na}^+]$ (into the fluid) and $[\text{Ca}_{5-y}, \text{REE}_y][\text{P}_{3-y}, \text{Si}_y]\text{O}_{12}\text{F} + y [\text{Ca}^{2+} + 2 \text{P}^{5+}]$ (from the fluid) = $\text{Ca}_5\text{P}_3\text{O}_{12}\text{F} + y \text{REEPO}_4 + y [\text{Si}^{4+}]$ (into the fluid)

(Harlov 2015). The final composition of the (REE + Y)-phosphate inclusions depends on the chemistry of the altering fluid, the P–T conditions of the alteration and the host apatite chemistry (Harlov and Förster 2003). Therefore, the presence of different (REE + Y)-phosphate inclusions in different zones of the host apatite can be explained by differences in the Y and REE compositions. Furthermore, Birski et al. (2018) concluded that the REE-F complexes in an H_2O solution and the specific budgets of REEs and Y in the apatite were responsible for the nucleation and formation of the (Ca–Y)-phosphate phases in the host apatites in the rocks of the BGB. Finally, in both samples, an orientation relationship between the solid inclusions and the host apatite is observed. In AL03-29B, an epitaxial relationship between the $\text{Ca}_3\text{Y}(\text{PO}_4)_3$ inclusions and the host apatite has been shown (Birski et al. 2018), whereas in sample H30, an orientation relationship has been observed between the monazite inclusions and the apatite. Previous researchers (Pan et al. 1993; Pan and Fleet 2002; Harlov 2015) have noted that REE-phosphate inclusions, e.g., monazite, can be highly elongated. This elongation often occurs along the *b* axis and parallel to the *c* axis of the host apatite (Harlov 2015). This feature has been observed in the analyzed monazite inclusions (Fig. 8e), where the [010] direction of the monazite is always parallel to the [0001] direction of the apatite. According to Harlov et al. (2007), this phenomenon can be explained by a topotactic reaction, i.e., a chemical solid-state reaction in which the orientations of the product crystals are determined by the orientation of the initial crystal. In this case, the formation of monazite does not involve exsolution from a solid solution; instead, it involves the fluid-aided growth of a separate phase that utilizes the available chemical components. This growth can be referred to as topotactic because the formation and growth of one phase (i.e., monoclinic monazite) within a second phase (i.e., apatite) can be required to accommodate the differences in the lattice parameters and d-spacings. An orientation relationship is not visible in the xenotime inclusions. This may be explained by the differences in the crystal lattice systems among the $\text{Ca}_3\text{Y}(\text{PO}_4)_3$, $\text{CaYP}_7\text{O}_{20}$ (Birski et al. 2018), monazite and xenotime inclusions and the similarities in the lattice spacings between the host apatite and the monazite and $\text{Ca}_3\text{Y}(\text{PO}_4)_3$ inclusions. All these observations together suggest that the REE + Y-phosphate inclusions originated via dissolution–reprecipitation processes. The formation is not dated, and consequently there is no evidence as to whether these changes occurred within the Archaean hydrothermal or a later system.

Igneous apatite

The REE profile of population A in sample H30 indicates that, at least to a limited extent, a signature of magmatic

origin could be potentially preserved. According to previous studies (Lepland et al. 2002; Higashi et al. 2017; Wudarska et al. 2018), REEs are more resistant to metasomatic alteration than the volatile elements that substitute into the apatite structure. Therefore, the REE distribution patterns should more clearly demonstrate a magmatic origin. A very distinguished negative Eu anomaly together with a slight enrichment in LREEs seems to confirm late crystallization from a differentiated melt. The yttrium behavior in apatite is similar to that of REEs, especially holmium, which has an identical valence and similar ionic radii. Therefore, Y and Ho behave similarly in many geochemical processes, resulting in a constant chondritic weight ratio $Y/Ho = 28$ in igneous rocks (Bau et al. 1996) and in hydrothermal fluids (Bau and Dulski 1999). Hence, the Y–Ho ratios do not indicate whether the apatite is of magmatic or metasomatic origin (Fig. 10). The features observed in CL suggest a complex history. A blue–violet core is characteristic of REE^{3+} emission centers. Alkaline rocks have been proven to tend towards blue, violet or lilac CL colors (Kempe and Götze 2002). In contrast, Mn^{2+} emission centers become dominant over bands characteristic of REEs in areas that are closer to the rims. This observation confirms the progressive alteration and transformation of the initially magmatic apatite. Since orange luminescence (activated by Mn^{2+}) is typical for acidic environments (Kempe and Götze 2002), the fluids that metasomatically altered apatite crystals might be related to the granitic magmatism at 3.45, 3.22, and 3.1 Ga (Kamo and Davis 1994; Dziggel et al. 2002). The acidic metasomatism of the volcani-sedimentary rocks deposited in the BGB has been confirmed (Rouchon and Orberger 2008). Moreover, differences in the Y and REE compositions are clearly visible in CL images (Fig. 8c) in the form of more blue and violet zones that might reflect variations in the REE and Y budgets of the magma during the crystallization of apatite. TEM observations seem to confirm the partial preservation of the igneous signature (Fig. 8h). The obtained results allow the assignment of the apatite from this group to the Archaean environment with the preservation of many of its original features.

Conclusions

The growth/regrowth morphology patterns, compositional variations, and alterations of the studied Archaean apatite provide valuable information on the environmental conditions during their formation and transformation. Based on that, three generations of apatite occurring in the BGB can be designated as follows:

A. Early-generation apatite that precipitated from Archaean seawater. This group clearly preserved the features that

can be expected from apatite precipitating before the GOE. This phosphate is Cl-depleted fluorapatite with a significant amount of OH. The REE distribution patterns are flat with positive Eu anomalies and without a negative Ce anomaly, and the preserved Y–Ho ratios of 54–70 are comparable to those of modern seawater (Bau and Dulski 1994). The CL data indicate precipitation under alkaline conditions. A characteristic of the apatite in this group is the presence of epitactically intergrown pyrrhotite inclusions. TEM investigation does not reveal any indicators of secondary alteration processes in this apatite, and the FTIR spectra indicate that the sedimentary apatites are relatively rich in water. Mn and Sr substitutions cause a stretching mode in the hydrogen bonding between OH and F and the consequent presence of additional component bands.

- B. Late-generation fluorapatite that experienced metasomatism/metamorphism. Their secondary origin is confirmed by REE distribution patterns depleted in both LREEs and HREEs. CL data suggest an origin in acidic metasomatism, most likely associated with granitic magmatism at 3.45–3.1 Ga (Kamo and Davis 1994; Dziggel et al. 2002). TEM observations reveal the presence of many structures such as cracks, pores and nanochannel-like structures as well as the occurrence of (REE + Y)-phosphate inclusions. FTIR measurements confirm that the metasomatic apatites are OH-group depleted. Moreover, the removal of trace elements that substitute in the Ca2 position from the structure of apatite causes a lack of any additional band present in the early-generation apatites of the first group.
- C. Igneous-derived apatite that partially preserves its igneous origin. These grains are predominantly fluorapatite such as those in the second group; however, their REE distribution patterns are slightly enriched in LREEs and slightly depleted in HREEs and have distinct negative Eu anomalies. The CL data indicate that the apatites in this group preserve the signature of igneous origin within their cores, whereas the rims are metasomatically altered. Due to the predominant F substitution in the X-site, the OH-stretching mode is the strongest for the OH_F band. Moreover, the occupancy by Mn and Sr of the Ca2 position is visible in the FTIR spectrum as additional $Mn-OH_F$ and $Sr-OH_F$ bands.

It seems that some of the apatite partly preserved its primordial features and therefore might be used for the reconstruction of Archaean abiotic systems. In this manner, some kind of comparison can be made between the periods before and after the GOE.

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