



Research Paper

Nanorocks, volatiles and plate tectonics

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ABSTRACT

The global geological volatile cycle (H, C, N) plays an important role in the long term self-regulation of the Earth system. However, the complex interaction between its deep, solid Earth components (i.e. crust and mantle), Earth's fluid envelopes (i.e. atmosphere and hydrosphere) and plate tectonic processes is a subject of ongoing debate. In this study we want to draw attention to how the presence of primary melt (MI) and fluid (FI) inclusions in high-grade metamorphic minerals could help constrain the crustal component of the volatile cycle. To that end, we review the distribution of MI and FI throughout Earth's history, from ca. 3.0 Ga ago up to the present day. We argue that the lower crust might constitute an important, long-term, volatile storage unit, capable to influence the composition of the surface envelopes through the mean of weathering, crustal thickening, partial melting and crustal assimilation during volcanic activity. Combined with thermodynamic modelling, our compilation indicates that periods of well-established plate tectonic regimes at <0.85 Ga and 1.7–2.1 Ga, might be more prone to the reworking of supracrustal lithologies and the storage of volatiles in the lower crust. Such hypothesis has implication beyond the scope of metamorphic petrology as it potentially links geodynamic mechanisms to habitable surface conditions. MI and FI in metamorphic crustal rocks then represent an invaluable archive to assess and quantify the co-joint evolution of plate tectonics and Earth's external processes.

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1. Introduction

The global volatile cycle (e.g. H, C, N), key to the long-term, self-regulation of the Earth system, relies on the continuous redistribution of elements between Earth's fluid envelopes (i.e. atmosphere and hydrosphere) and the its solid layers (i.e. crust and mantle) (Houlton et al., 2018; Lee et al., 2019). Introduction of volatiles into the terrestrial mantle may have triggered plate tectonics, which as best as we know could have been important for the emergence of complex life (Albarede, 2009). In the Phanerozoic, volcanic activity and subduction drive volatile exchanges between the solid Earth and the surface in extensional and convergent settings (e.g. Kelemen and Manning, 2015; Lee et al., 2019; Plank and Manning, 2019; Wong et al., 2019; Werner et al., 2019). This has been used to support the idea that plate tectonics, through the expression of the Wilson cycle (subduction – collision – rifting) might control in turn the long-term volatile cycle (e.g. Wong et al., 2019 and references therein). The stability of the Earth system would then depend on the positive feedback between surface

conditions and geodynamic mechanisms (Lenardic et al., 2019; Sobolev and Brown, 2019).

In collisional settings, reworking of the continental crust is the result of the combination of partial melting, deformation and melt segregation processes. The partial melting of buried supracrustal lithologies (e.g. Chappell and Stephens, 1988; Clemens, 1990, 2003; Villaros et al., 2009; Clemens et al., 2011) mobilises a substantial amount of mineral-bound volatiles (e.g. H₂O, CO₂) at deep crustal levels (e.g. Nicoli and Dyck, 2018; Nicoli, 2019). Along the PT path, the formation of anatectic liquids ultimately leads to the formation of S-type granites, after extraction from the source region and migration into the shallow crust (Clemens et al., 2011). The cogenetic links between the source, the migmatites and the resulting granites (e.g. Nicoli et al., 2017; Nicoli and Dyck, 2018; Clemens, 1990), embodies directly the connection between erosion and weathering processes at the surface, responsible for the formation, the volume and the composition of sediments (Condie, 1993), and the metamorphism and melting at depth. The capacity of the surface reservoir signature (i.e. organic N and C isotopic signature) to be retained beyond partial melting conditions in the lower crust indicates that a significant amount of volatiles can be isolated from the surface cycle and stored in

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Table 1

List of case studies of primary melt inclusions (both nanorocks and glassy) and/or fluid inclusions in peak minerals from crustal rocks. When fluid presence during melting is inferred, i.e. not directly verified via identification of primary fluid inclusions, this is reported in the "Note" column. The criteria used to compile this dataset are discussed in the text.

Locality	Age (Ma)	$\pm 1\sigma$	P (GPa)	$\pm 1\sigma$	T (°C)	$\pm 1\sigma$	T/P	Rock	Protolith	Host	MI	FI	Fluid	Note on inclusions	Ref.
El Hoyazo - Mazarrón (Spain)	9.6	0.3	0.60	0.1	850	50	1417	Granulitic enclaves	Sediments	Grt/Pl/Crd/Spl	x	x	COH-N		Acosta-Vigil et al. (2007); Cesare et al. (2007); Ferrero et al. (2011)
Seram, eastern Indonesia	16.0		0.80		950		1188	Felsic granulites	Sediments	Garnet	x				Pownall et al. (2019)
Barun Gneiss (Himalayas)	30.0	8	0.80		830	30	1038	Migmatites	Sediments	Garnet	x				Ferrero et al. (2012)
Gruf granulite	33.0	4	0.90	0.05	900	10	1033	Felsic granulites	Sediments	Garnet	x	x	COHN	C-Dominated inclusion	Gianola et al. (2020)
Kali Gandaki (Himalayas)	38.5	2.5	1.05	0.05	685	35	652	Migmatites	Sediments	Garnet	x				Bartoli et al. (2019); Carosi et al. (2015)
Rhodope Metamorphic Province (Greece)	40.0		1.35	0.15	775	25	574	Paragneisses	Sediments	Garnet	x			Carbonate inclusions	Mposkos et al. (2009)
Maghrebian basement (Tunisia)	12.0	2	0.55	0.15	780	20	1418	Xenocrysts	Granitoids	Garnet	x	x	COH-N		Ferrero et al. (2014)
Altai orogenic belt (China)	255.8	1.8	0.45	0.05	850	50	1889	Mafic granulites	Mafic rocks	Garnet		x	Pure CO ₂	C-Dominated inclusion	Li et al. (2004)
Jubrique, Betic Cordillera (Spain)	287.0	4	1.30	0.1	850		654	Migmatites	Sediments	Garnet	x	x	CO ₂ -rich		Barich et al. (2014)
Ivrea Zone (NW Italy)	270.0	30	0.80		800		1000	Migmatites	Sediments	Garnet	x	x	COH-N		Carvalho et al. (2019)
Altai orogenic belt (China)	284.1	2.6	0.80		980		1225	Pelitic granulites	Sediments	Garnet	x				Liu et al. (2020)
Ronda migmatite (Spain)	286.0	11	0.48	0.03	680	20	1432	Migmatites	Sediments	Garnet	x				Bartoli et al. (2014)
Oberpfalz area, SW Bohemian Massif (Central Europe)	325.0	50	0.65	0.15	800		1100	Migmatites	Sediments	Garnet	x	x	COH-N	Nanocarbonatites	Ferrero et al. (2016a)
Bavarian Unit (Bohemian Massif)	335.0	5	0.60		870	40	1450	Paragneisses	Sediments	Garnet	x				Ferrero et al. (2018b)
Blansky Lés (Bohemian Massif)	335.0	5	2.25	0.50	950		422	Felsic granulites	Granitoids	Garnet	x				Ferrero et al. (2018b)
Hartsteinwerk loja quarry (Bohemian Massif)	335.0	5	0.75	0.15	775	25	1033	Paragneisses	Sediments	Garnet	x				Ferrero et al. (2018b); Sorger pers.comm.
Ktis (Bohemian Massif)	335.0	5	1.80	0.4	800	100	889	Paragneisses	Sediments	Garnet	x	x	COH-N		Ferrero et al. (2018b)
Mohsdorf (Bohemian Massif)	335.0	5	1.60		1000		625	Felsic granulites	Granitoids	Garnet	x				Ferrero et al. (2018b)
Plaimberg (Bohemian Massif)	335.0	5	4.50	0.50	1000	100	244	UHP eclogites	Mafic rocks	Garnet	x				Ferrero et al. (2018b)
Saidenbach (Bohemian Massif)	335.0	5	4.50	0.50	1000		222	Felsic granulites	Sediments	Garnet	x			Diamond	Stöckhert et al. (2001)
Steinaweg (Bohemian Massif)	335.0	5	1.85	0.45	825	25	446	Felsic granulites	Granitoids	Garnet	x				Ferrero et al. (2018b)
Stuckestein (Bohemian Massif)	335.0	5	0.60	0.10	800		1333	Migmatites	Sediments	Garnet	x	x	COH-N		Ferrero et al., (2018b)
T7 borehole (Bohemian Massif)	335.0	5	4.50	0.50	1000		222	Felsic granulites	Sediments	Garnet	x			Diamond	Kotkova et al. (2014)
Erzgebirge (Bohemian Massif)	340		4.5		1000		222	Eclogites	Basaltic dyke?	Garnet	x			Metasomatic melt	Borghini et al. (in prep)
Granulitgebirge (Germany)	340.0		2.10	0.10	1000	7.5	480	Eclogites	Gabbro?	Garnet	x			Metasomatic melt	Borghini et al. (2020)
Orlica-Śnieżnik Dome (Bohemian Massif)	340.0	50	2.70		875	12.5	329	Felsic granulites	Granitoids	Garnet	x				Ferrero et al. (2015)
Central Maine terrane (USA)	379.0		1.80		1050		1000	Felsic granulites	Sediments	Garnet	x				Axler and Ague (2015); Ferrero et al. (2021)
Gory Sowie (Bohemian Massif)	400.0		0.75	0.10	950	50	1267	Felsic granulites	Granitoids	Garnet	x				Stupski et al. (2018)
Khabarny mafic-ultramafic Massif (Urals)	415.0	8	0.80	0.03	785	45	981	Garnetites	Mafic rocks	Garnet		x	COH-N		Bakker et al. (2020)
Seve Nappe Complex (Scandinavian Caledonides)	455.0	5	4.00		835	5	209	Paragneisses	Sediments	Garnet	x	x	COH-N	Diamond	Klonowska et al. (2017)
Mont Albert ophiolitic complex	456.0	5	1.00		850		850	Metapelites	Sediments	Garnet	x				Dubacq et al. (2019)
MT. Edixon (Antartica)	515.0	15	0.80	0.1	760	20	950	Schists	Sediments	Garnet	x	x	COH-N		Ferri et al. (2020)
Khondalite Belt (Southern India)	525.0	5	0.60	0.10	900		1500	Migmatites	Sediments	Garnet	x				Ferrero et al. (2012)
Kokchetav Massif (Kazakhstan)	530.0		4.75	0.25	975	25	205	Paragneisses	Sediments	Garnet	x			H ₂ O-K fluid	Stepanov et al. (2016)
Dronning Maud Land (Antartica)	542.0	27	1.55	0.15	910	50	587	Mafic granulites	Mafic dikes	Garnet	x				Ferrero et al. (2018b)
Hakuritale granulites (Sri Lanka)	550.0	50	0.95	0.05	830			Al, Mg-rich granulite	Metabasite?	Garnet		x	CO ₂ -rich		Bolder-Schrijver et al. (2000)
Lützow-Holm Complex, East Antarctica	550.0	50	0.95	0.25	875	75	921	Mafic granulites	?	Garnet	x				Saitoh and Tsunogae (2015)
Dronning Maud Land (Antartica)	565.0	15	0.69	0.04	1080		1565	Anorthosite	Anorthosite	Plagioclase		x	H ₂ O-	C-Dominated	Kleinefeld and Bakker

morphs, Ferrero and Angel, 2018) along with OH-bearing minerals and, in few cases, limited amounts of CO₂ and H₂O in interstitial position (Bartoli et al., 2013) (Fig. 1b). Consistently with these observations, such inclusions were originally named *nanogranites* (Cesare et al., 2009) and later *nanogranitoids* (Bartoli et al., 2016). Recent works however showed that crystallized melt inclusions in metamorphic rocks have a much larger variety in composition, i.e. from silicic to carbonatic, prompting Bartoli and Cesare (2020) to spearhead the use of the term “nanorock” as a more comprehensive and appropriate name, and, for this reason, this term will be used throughout the present work. Later studies (Carvalho et al., 2020; Ferrero et al., 2018a; Ferri et al., 2020) have then demonstrated that, besides the anatectic melt, also the fluids present during partial melting can be trapped as primary inclusions in migmatites (Fig. 1c). These fluids consist of CO₂, CH₄, N₂ and H₂O in variable proportions (e.g. Carvalho et al., 2020; Ferrero et al., 2018b; Tacchetto et al., 2019) (Fig. 1d).

Importantly, melt inclusions in metamorphic garnet do not suffer issues commonly recognised in magmatic melt inclusions, such as volatile loss and doubtful correspondence between trapped and original melt due to compositional heterogeneities present in the melt immediately adjacent the growing crystal (i.e. boundary layer effect – see discussion in Cesare et al., 2015). In particular, while diffusive H₂O and CO₂ loss is common for magmatic melt in olivine (Gaetani et al., 2012), driven by the pressure gradient between trapped melt and surrounding magma under continuous depressurisation during ascent, such a driving force is absent in rock-dominated system (i.e. metamorphic rocks during cooling – Bartoli et al., 2014). The H₂O and CO₂ content measured in crack-free melt inclusions in metamorphic garnets can be thus considered as representative of the original volatile content of the melt at depth (more details on the topic in Bartoli et al., 2014; Cesare et al., 2015; Ferrero and Angel, 2018). Boundary layer effects only affect trace elements highly enriched in garnet (e.g. Y, HREE, Acosta-Vigil et al., 2012), thus making the anatectic inclusions overall representative of the original melt composition.

A growing body of evidence (e.g. Carvalho et al., 2020; Ferrero et al., 2016a; Tacchetto et al., 2019 and references therein) has revealed a significant variation in the nature, abundance and distribution of fluid-bearing inclusions in metamorphic minerals. Such variations can be interpreted as resulting from differences in the chemistry of the source, fluid percolation prior entrapment, as well as specific geodynamic settings, which will influence the dehydration/decarbonation sequences along the metamorphic prograde path (e.g. Nicoli and Dyck, 2018; Stewart and Ague, 2018; Stewart et al., 2019 – see section 4 below). The quantification of immiscible pristine COH(±N) fluid phases in rehomogenised nanorocks using NanoSIMS (e.g. Bartoli et al., 2014; Carvalho et al., 2020; Ferri et al., 2020; Ferrero et al., 2021) together with thermodynamic modelling is increasingly becoming a reliable approach to quantify reaction kinetic and thermodynamic equilibrium in the deep continental crust. Importantly, all these applications reassessed the critical role played by fluid-assisted partial melting in accretionary settings, previously relatively neglected – apart from a few remarkable exceptions (e.g. Sawyer, 2010; Weinberg and Hasalová, 2015).

3. Inclusions in space and time

In the present work we employed a database (Table 1) of 46 localities where primary melt (MI) and/or fluid inclusions (FI) have been reported in minerals formed during prograde and peak regional metamorphism of crustal rocks. Out of the whole database, thirty case studies contain only MI, six case studies only FI and ten cases a combination of both trapped at the same time (i.e.,

under primary fluid-melt immiscibility conditions; see Cesare et al., 2015). The term “MI” refers to polycrystalline inclusions originally trapped as a melt (regardless of their composition, i.e. either silicate rich or carbonatic), partially crystallized inclusions as well as glassy inclusions. For the purpose of our study, such inclusions are to be considered informative of fluid/melt regime at depth because of their primary nature, i.e. they formed during the growth of the host phase (Roedder, 1984), as well as of the absence of evidence for decrepitation. Fluid inclusions present a different caveat: as thoroughly discussed by Carvalho et al. (2020), most (all?) of the COH(±N) fluid inclusions found in garnet underwent post entrapment changes on cooling with crystallization of new phases. Although this largely prevents a precise quantification of the original fluid composition, their characterisation provides nevertheless precious data on fluid speciation at depth.

Geographically, these findings span all continents with the exception of South America and Australia, and the inferred protolith of the host rocks cover the whole range of rocks, from sediments to granitoid to mafic rocks, involved in continent–continent collision. The most common host for inclusions is peritectic garnet, which is hardly surprising in view of its abundance in metamorphic rocks of crustal origin and its vast range of stability and refractory nature. In some cases, peritectic plagioclase (Madlakana and Stevens, 2018; Nicoli et al., 2017) may also be an important host of inclusions (three case studies). Both peritectic garnet and peritectic plagioclase form in the presence of melt and are cogenetic with MI. The inspection of the database of MI in metamorphic rocks reported by Cesare et al. (2015) shows that zircon is the second most common inclusion host in crustal rocks. However, findings of MI in zircons are left out of the present compilation: zircon crystallizes more often on cooling during the retrograde evolution of the rocks (Roberts and Finger, 1997), and in absence of detailed studies on each case, it is impossible to relate with certainty MI in zircons to the prograde/peak history of the host rocks.

The inclusions discussed here sampled portions of melt and/or fluids present in the deep crust between ~680 °C (Ronda migmatites) and 1050 °C (Central Maine Terrane felsic granulites), and 0.45 GPa (Altai orogenic Belt) to 4.75 GPa (subduction of the continental crust in Kokchetav Massif). Most observations occur at 750–900 °C and ~0.5–1.5 GPa, which correspond to the stability field of granulitic rocks (Harley, 1989) (Fig. 2a). The limited number of findings of MI and FI in ultrahigh pressure (UHP) and ultrahigh temperature (UHT) crustal rocks may be simply related to the relative rarity of such rock types, whereas their absence in high-grade – low pressure crustal rocks is likely due to the absence of peritectic garnet, the most common and reliable host of melt/fluid inclusions, often replaced by the much less robust cordierite. Whereas no particular correlation is visible between different groups and particular PT conditions and/or PT gradients, evidence for the coexistence of crustal melt and a fluid are concentrated in the temperature range 780–900 °C. The fluid is mainly COH(±N) in composition, with a high CO₂ component in most cases (e.g. Carvalho et al., 2019; Cesare et al., 2007; Ferrero et al., 2014, 2018a; Gianola et al., 2020), only two cases showing a H₂O-bearing fluids (Ferrero et al., 2011; Ferri et al., 2020).

For the purpose of this study, the assessment of the role of continental crust evolution in the stabilisation for the global volatile cycle, the most crucial aspect of these findings is their distribution in time (Fig. 2b). The list of case studies covers most of the Earth's history, from 2.8 Ga (Uttental, Greenland) to 9.6 Ma (El Hoyazo, Southern Spain). Most of these inclusions are found in metasedimentary lithologies and few in granitoids and mafic rocks (Fig. 1b). Nanorocks and primary COH(±N) fluid inclusions are mainly a Phanerozoic affair (Fig. 1b). A second cluster of inclusions occurs between 2.5 Ga and 3.0 Ga. These two main periods also correspond to the formation of economically viable graphite

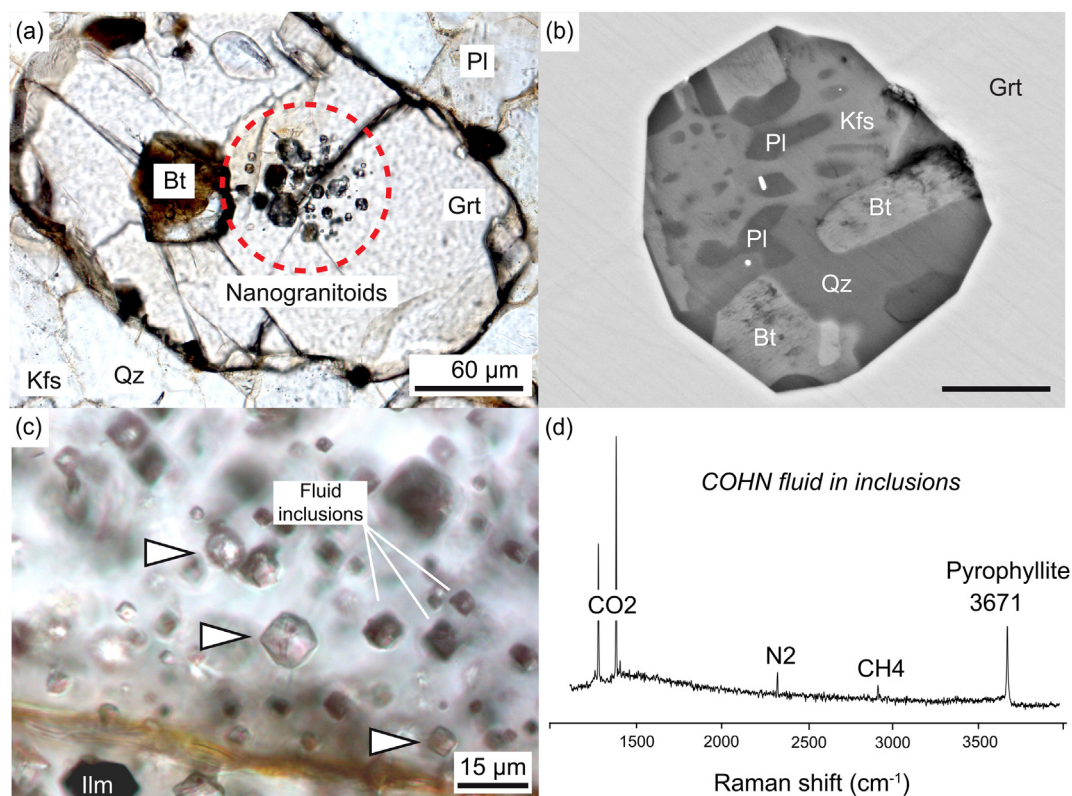


Fig. 1. Nanorocks and fluid inclusions in metamorphic crustal rocks. (a) Typical appearance of a cluster of nanogranitoids in garnet (Ferrero et al., 2016b). Kfs = K-feldspar, Qz = Quartz, Grt = garnet, Pl = plagioclase, Bt = biotite. (b) Nanogranitoid from Koliakkode quarry, Kerala Khondalite Belt (Southern India; PhD thesis S. Ferrero). (c) Nanogranitoids (white arrows) and fluid inclusions in metasediments trapped under conditions of primary fluid-melt immiscibility. Ilm = ilmenite. (d) Typical Raman spectrum of fluid inclusions coexistent with nanogranitoids (Carvalho et al., 2020). Both (c) and (d) are modified after Ferrero et al. (2016a).

deposits (Luque et al., 2014), arguing for major carbon-bearing fluid fluxes in the lower crust (Touret et al., 2016). S-type granites (e.g. Clemens et al., 2011; Zhu et al., 2020) show the same distribution pattern as sediment related inclusions, with a third important peak during the assembly of the Columbia-Nuna supercontinent (1.7–2.1 Ga) (Fig. 2c). Whereas there is evidence of intrusions during the Boring Billion (0.85–1.7 Ga) (Roberts, 2013), only one inclusion in metasedimentary rocks has been identified within this period so far. The issue of preservation bias is mentioned in the discussion below.

The widespread occurrence of inclusion findings across a large range of PT conditions, different protoliths and ages show the capability of natural rocks to sample and preserve deep crustal fluids, making these inclusions reliable probes to investigate planetary scale processes such as fluid regime variations in relationship with plate tectonics. The joint occurrence of nanorocks in metasedimentary rocks, graphite deposits and crustal derived magmas at specific time of Earth's history (Fig. 2b,c) suggests that there might be periods prone to enhanced coupling between crustal reworking and surface mechanisms.

4. Protolith chemistry and partial melting conditions

Nanorocks represent melts and fluids in equilibrium with the bulk rock residue (Acosta-Vigil et al., 2017). The composition of primary melt and fluid phases depends on the chemistry of the source material (e.g. Le Breton and Thompson, 1988; Clemens and Stevens, 2012). In the continental crust, minor compositional variations (e.g. Fe + Mg and K contents) (Condie, 1993) in the upper crust can lead to significant variations in the depth and sequence of devolatilisation events and influence the nature of partial melting

reactions and the chemistry of the resulting magmas. For instance, changes in the composition of siliciclastic and volcanoclastic sediments from the Archean to the Phanerozoic are reflected in the evolution of anatectic liquid chemistry towards an increasingly Mg-poor, Ca-rich, less viscous monzogranitic melt (Nicoli and Dyck, 2018; Nicoli, 2019). Although data on the volatile content of primary melt inclusions is still scarce, we compared H₂O and CO₂ contents of rehomogenised inclusions with proposed protolith compositions (Fig. 3a). Protolith with high Ca/(Ca + Na) and Mg# ($Mg\# = Mg^{2+}/(Mg^{2+} + Fe^{2+})$) and low-K tend to favour anatectic melt with higher H₂O and CO₂ contents. This is consistent with previous findings (Nicoli and Dyck, 2018; Nicoli, 2019) which have showed that the ferromagnesium content of siliciclastic rocks influences the temperature of mica breakdown and the amount of water available at solidus conditions. Calc-silicate, meta-carbonate, and high-Ca siliciclastic rocks are able to retain greatest quantities of carbon under metamorphic conditions typical of accretionary and collision orogenies. This is because most of the metamorphic carbon is held in the crystal structure of the carbonate minerals, calcite [CaCO₃] and dolomite [CaMg(CO₃)₂], and the stability of these minerals is greatest in compositions high in CaO, SiO₂, and MgO. The composition of primary melt and fluid phases is also determined by where and when along the prograde path they are produced and then entrapped into growing peritectic phases, which then relates to the degree of partial melting. At the mineral scale, the composition is influenced by the interplay between kinetics of diffusion between melt and minerals, the possibility of mineral recrystallization and the time gap between the onset of partial melting and entrapment (Acosta-Vigil et al., 2017). Therefore, the nanorock archive and associated fluid phases in metasedimentary rocks have been primarily used to investigate

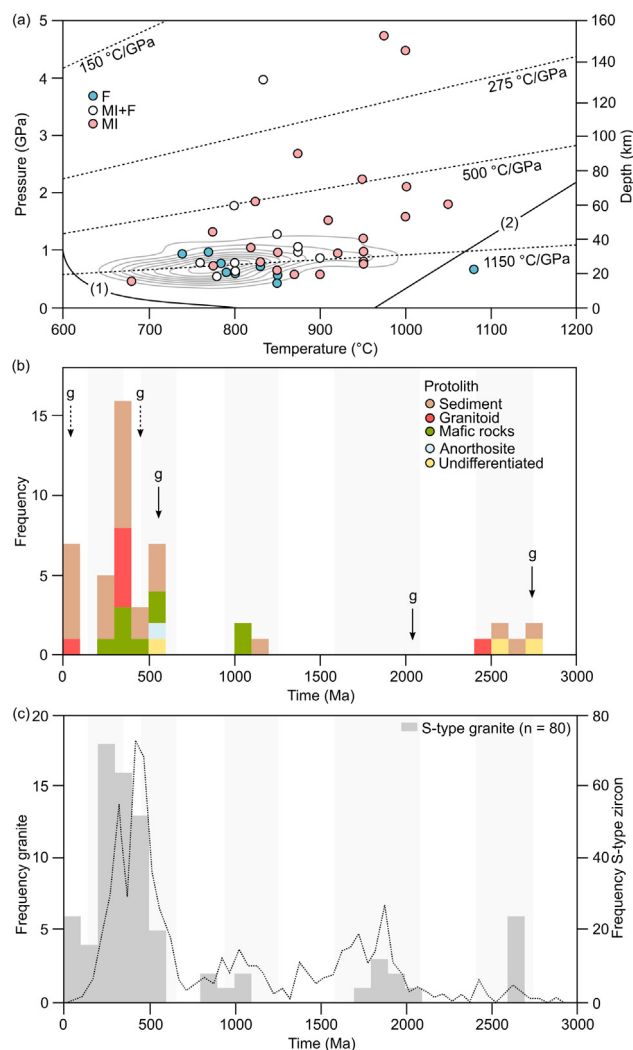


Fig. 2. Nanorocks and primary COH(±N) fluid inclusions. (a) Metamorphic conditions at which melt and fluid inclusions are entrapped by growing peritectic mineral phases (e.g. garnet, plagioclase). 1 - wet granite solidus; 2 - dry granite solidus. The kernel density represents the domain of equilibration of granulitic rocks, from 3.0 Ga to the present (Harley, 1989). (b) Main occurrence of inclusions in high-grade metamorphic mineral in different protoliths. Arrows represent principal graphite deposits (Luque et al., 2014). Dashed line: igneous deposit; plain line: metamorphic deposit. (c) Occurrence of S-type granites (updated database from Clemens et al., 2011; Taylor et al., 2014). The dashed line indicates the occurrence of S-type zircons reconstructed from river sediments (Zhu et al., 2020). Vertical shaded areas represent supercontinent assembly phases.

the primary composition of anatectic melts (e.g. Cesare et al., 2011, 2015; Bartoli et al., 2014), the nature of the melting and reaction and fluid budget of the continental crust (e.g. Acosta-Vigil et al., 2010; Carvalho et al., 2019; Ferrero et al., 2011, 2014, 2016a; Ferri et al., 2020; Gianola et al., 2020; Tacchetti et al., 2019; Acosta-Vigilet al., 2007).

To investigate further the relation between the H₂O and CO₂ content of nanorocks and secular changes in the composition and the thermal state of the crust, we used phase equilibria modelling on average shale bulk compositions from the Archean (2.5–3.0 Ga), Proterozoic (0.6–2.5 Ga) and Phanerozoic (0–0.6 Ga) (Condie, 1993). Shales, primary source of anatectic melts, form from the accumulation of products of weathered continental material in passive continental margins, later remobilised during continental collision (Condie, 1993; Bradley, 2008). We then defined a set of PT paths by considering minimum and maximum orogenic metamorphic gradient (i.e. T/P > 500 °C/GPa⁻¹) over periods of 200 Ma

(Brown and Johnson, 2018). We used version 6.9.0 of the Perple_X software package (Connolly, 2009) and the 2004 update of the Holland and Powell (2011) thermodynamic database (<http://www.perple.ethz.ch/>) in the system Na₂O-CaO-K₂O-FeO-MgO-Al₂O₃-SiO₂-H₂O-TiO₂ (NCKFMASHT). The solution models used are as follows: silicate melt model, garnet, chlorite, muscovite, biotite, orthopyroxene after White et al. (2014); feldspar (Fuhrman and Lindsley, 1988); amphibole model after Diener and Powell (2012). First, we followed the same approach used by Nicoli and Dyck (2018) and individually fixed bulk-rock water content at the point at which the relevant geotherm intersects the solidus to permit water-absent partial melting. Results are presented in Table 2 and Fig. 3b.

Nanorocks contain a variable amount of H₂O, from 2 wt.% to 14 wt.% (Bartoli et al., 2014, 2019; Carvalho et al., 2019; Ferri et al., 2020) (Fig. 3a). Our modelling reproduces the entire range of observed H₂O contents in anatectic inclusions. The secular evolution of the thermal state of the crust (i.e. apparent metamorphic gradient, T/P - Brown and Johnson, 2018), combined with changes in the shale chemistry (Condie, 1993) directly influences the water content of the melt (Nicoli and Dyck, 2018). Our modelling shows that maximum water contents in the melt is reached for low T/P (<1000 °C/GPa⁻¹) gradient at 0–1.2 Ga, 1.8–2.0 Ga and 2.4–2.8 Ga, corresponding to periods of well-established plate tectonic mechanisms, generalisation of mobile lid tectonics and transition from stagnant lid to mobile lid tectonics respectively (e.g. Condie, 2018; Holder et al., 2019; Palin et al., 2020) (Fig. 3b). Minimum melt water contents are reached during the tectomagmatic lull at 2.2–2.3 Ga (Spencer et al., 2018) and the Boring Billion, during which single lid tectonic regimes might have been the dominant geodynamic regimedominated (Stern, 2020). Our modelling also suggests that the range of melt water content for high T/P (>1000 °C/GPa⁻¹) might have been constant since the Archean.

In order to estimate the concentration of CO₂ in the combined melt and the fluid phase, we used the Phanerozoic shale composition in the NCKFMASHT-CO₂, adding a fluid phase (Connolly and Trommsdorff, 1991) and carbonates (Holland and Powell, 2011; Kuhn, 2004) to the solution model. We calculated X_{CO2} (X_{CO2} = CO₂/(CO₂ + H₂O)) between 750 °C and 1150 °C along the two metamorphic gradients encompassing the conditions of rehomogenisations, 650 °C/GPa⁻¹ and 850 °C/GPa⁻¹, at fixed bulk H₂O content, 1.75 wt.% and 1.35 wt.% respectively (Table 1) (Fig. 3c). The CO₂ content of nanorocks varies between 180 ppm and 8100 ppm (e.g. Carvalho et al., 2019; Ferri et al., 2020; Gianola et al., 2020; Ferrero et al., 2021). Although the data set is restrained to the Phanerozoic and Neoproterozoic, it is worth noting that there is an apparent positive correlation between concentration in CO₂ in the melt and the temperature (Fig. 3c). The modelling reproduces inclusions with CO₂ concentration <3000 ppm for a maximum X_{CO2} of 0.02. Higher CO₂ concentration values are reached for X_{CO2} between 0.05 and 0.1. It is worth noting that our modelling only considers a single source (i.e. shale). In reality, sediments in passive margins range from quartzite to volcanoclastic to carbonate (Veizer and Mackenzie, 2004). Hence, our modelling only provides an overview of the melting/volatile relationship in the deep continental crust.

5. Discussion

5.1. Connecting nanorocks to the global volatile cycle

The presence of CO₂ in the lower crust has been long argued to be the results of slab decarbonation and fluid flux from the mantle wedge since the late Archean (Newton et al., 1980; Newton, 1987; Santosh and Omori, 2008; Touret et al., 2016). Although ascending CO₂ percolation might still have occurred at the crust/mantle inter-

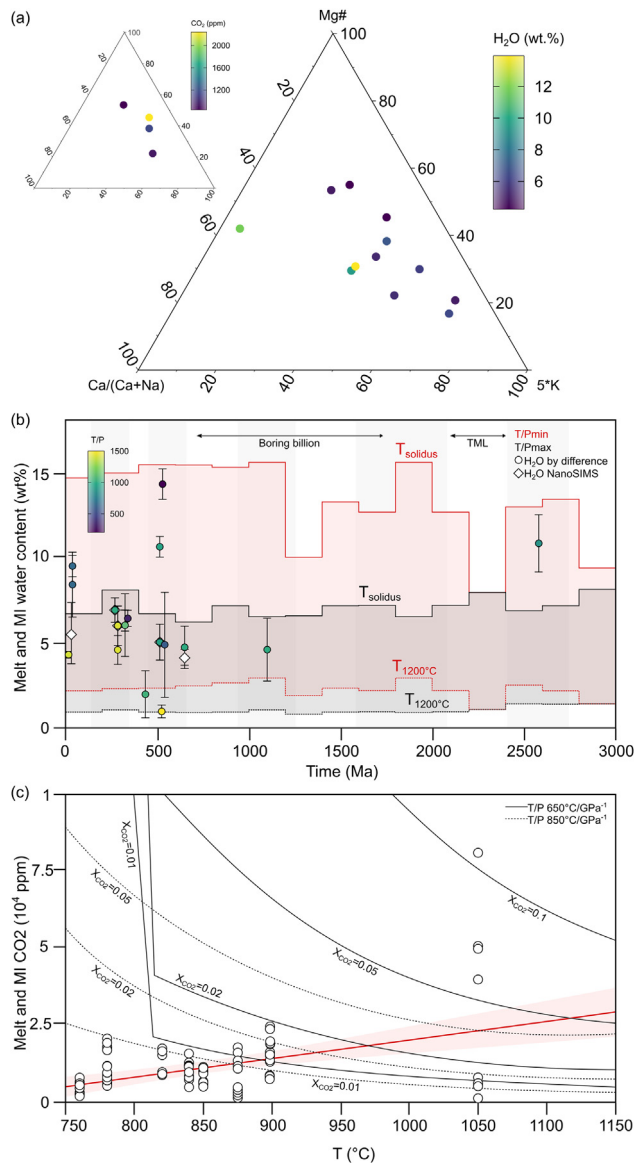


Fig. 3. Water and carbon dioxide content in rehomogenised melt inclusions from metasedimentary rocks estimated by difference from EMP totals or via NanoSIMS analyses. (a) H₂O and CO₂ content in inclusion plotted as a function of bulk rock protolith in Ca/(Ca + Na) – K – Mg# space. Mg# = $Mg^{2+}/(Mg^{2+} + Fe^{2+})$. Both ternary diagrams have identical axes. (b) H₂O content in modelled melt and melt inclusions (MI). The shaded areas represent the range of water content between T_{solidus} and T₁₂₀₀ for the minimum and maximum metamorphic gradients for each bin of 200 Ma. Vertical bars areas represent supercontinent assembly. TML: tectomagmatic lull (Spencer et al., 2018). The colour code for each datapoint correspond to the apparent metamorphic gradient (T/P). (c) CO₂ content in modelled melt and melt inclusions (MI) for different X_{CO₂} (i.e. CO₂/(H₂O + CO₂)) as a function of rehomogenisation temperatures. Linear regression on the average CO₂ content at given temperature with a 1σ error envelope (grey area) is also shown.

face, the primary nature of the nanorocks and associated primary COH(±N) fluids argues in favour of an internally derived source, making these inclusions preserved snapshots of deep crustal reworking processes. Therefore, their presence in metasedimentary rocks has a great potential as it offers a new tool to understand and quantify the transfer of volatile elements from the surface (e.g. formation of the protolith via weathering and erosion) to the lower crust.

In this part of the discussion, we show that the occurrence of MI and FI in high-grade rocks might be linked to the different phases

of the Wilson cycle during the Phanerozoic, which as best as we know controls the modern carbon cycle (Wong et al., 2019).

During continental break-up phases (50–150 Ma, 400–500 Ma; Condie et al., 2015), atmospheric CO₂ increased mainly as the results of crustal extension and degassing from rift zones (Berner and Kothavala, 2001; Foley and Fischer, 2017; Wong et al., 2019) (Fig. 4a). During continental assembly phases (150–400 Ma and >500 Ma), the increase in orogenic activity was followed by an increase in the weathering rate (François et al., 1993), drawing CO₂ from the atmosphere into siliciclastic sediments (Berner and Kothavala, 2001) (Fig. 4a). Continental silicate weathering is estimated to currently consume 38–75 MT C-yr⁻¹ (e.g., Gaillardet et al., 1999 and references therein). Carbon-bearing sediment amalgamate in passive margins, which are in turn remobilised during continental collision (Bradley, 2008). The burial of supracrustal rocks could then bring a substantial amount of CO₂ and H₂O towards crustal reworking sites. Along the way, devolatilisation reactions releases COH(±N) fluids into the middle crust. Steward et al. (2019) estimated that 2–23 MT C-yr⁻¹ are currently produced by metamorphic degassing in collision settings. If we consider these current outgassing rates and that up to half of the weathered continental silicate material deposited in passive margin is ultimately remobilised during crustal thickening, up to 50% of the carbon remains trapped in the root of orogenic belts and preserved passed partial melting conditions.

The occurrence of MI and FI in the geological record coincides with episodes of crustal thickening and high melt productivity (>50 vol%, Fig. 4b). These different observations highlight a ~320 Ma cyclicity for the transfer of volatiles from the crustal reservoir to the surface reservoir (Fig. 4b). This is in agreement with the tempo of orogenic activity (~80 Ma) (Brown et al., 2020), the 300 Ma periodicity of the magmatic/volcanic cycle (Brink, 2019) and the fact that the global flux of weathered material recycled into the mantle is currently balanced by addition of material at Earth's surface mainly in accretionary and collisional zones (Scholl and von Huene, 2007; Clift et al., 2009; Hawkesworth et al., 2010; Guo and Korenaga, 2020). Therefore, even if additional mechanisms need to be taken into consideration, such as glaciation (Foster et al., 2017) and cosmic cycles (Berger, 2013; Brink, 2019) (Fig. 4b), orogenic activity (weathering + metamorphism) might be capable to influence Earth's climate every ~160 Ma (Fig. 4b).

The continental crust is the third most important carbon reservoir on Earth (6.5×10^7 GT C), after the core (4×10^9 GT C) and the mantle (3×10^8 GT C) (DePaolo, 2015). Garnet was proposed to represent 4.9 vol% of the total crustal mass (Ronov and Yaroshevsky, 1969), and it is known to form mostly in the T range > 400 °C to > 1000 °C in metamorphosed crustal rocks (Baxter et al., 2013). As melt is stable from T > 700 °C, we considered that ~50% of the garnet forms in suprasolidus conditions (i.e. 2.5% of the total crustal mass). Assuming that 30–50% of the crystal volume of these garnets is occupied by inclusions containing 2–14 wt.% H₂O and 180 ppm and 8100 ppm CO₂ (Parisatto et al., 2018), the current maximum global amounts of water and carbon trapped in nanorocks are 10^5 – 10^6 GT and 10^3 – 10^5 GT respectively. As a comparison, these estimates would correspond to the equivalent of one Greenland ice sheet (2.3×10^6 GT H₂O) and one to two Earth's atmosphere (5.9×10^2 GT C) (Eakins and Sharman, 2010; Lee et al., 2019). These estimates are made assuming that all peritectic garnets have inclusions and all inclusions are preserved melt. Therefore, our mass balance calculation only gives an insight into the importance of the formation and stabilisation of a partially molten lower crust on the long-term storage of volatiles in the deep Earth (Touret, 2003). The presence of exposed late Archean metasedimentary granulitic rocks in the geological record (Harley, 1989) would then suggest the duration of this storage could easily exceed 2.5 Ga.

Table 2
Modelled bulk and melt water content along minimum (T/P_{min}) and maximum (T/P_{max}) orogenic metamorphic geothermal (T/P – Brown and Johnson, 2018) per bin of 200 Ma, using shale compositions from Condie (1993) as starting material.

Bin (Ma)	T/P _{min}					T/P _{max}				
	T/P (°C-GPa ⁻¹)	T _{solidus} (°C)	H ₂ O _{bulk} (wt.%)	T _{solidus} (wt.%)	H ₂ O _{melt} T ₁₂₀₀ (wt.%)	T/P (°C-GPa ⁻¹)	T _{solidus} (°C)	H ₂ O _{bulk} (wt.%)	T _{solidus} (wt.%)	H ₂ O _{melt} T ₁₂₀₀ (wt.%)
0–200	547	722	1.75	2.14	14.49	1500	722	0.88	0.91	6.71
200–400	518	722	1.75	2.24	14.92	1321	722	0.95	0.98	8.02
400–600	500	722	1.75	2.34	15.38	1500	722	0.88	0.91	6.71
600–800	507	722	1.75	2.34	15.38	1727	722	0.83	0.86	6.19
800–1000	571	722	1.7	2.60	15.18	1517	722	0.93	0.98	7.21
1000–1200	503	722	1.7	2.86	15.53	1629	722	0.93	0.98	6.53
1200–1400	850	735	1.53	1.82	9.91	2111	722	0.8	0.85	6.53
1400–1600	618	735	1.7	2.32	13.31	1500	722	0.93	0.98	7.21
1600–1800	682	735	1.7	2.25	12.61	1493	722	0.93	0.98	7.21
1800–2000	500	722	1.7	2.86	15.53	2000	722	0.8	0.85	6.53
2000–2200	650	735	1.7	2.25	12.61	1600	722	0.93	0.98	7.21
2200–2400	1269	722	0.93	1.00	7.92	1282	722	0.93	1.00	7.92
2400–2600	512	771	1.33	2.59	12.92	1313	735	1.45	1.52	6.92
2600–2800	621	735	1.36	2.25	13.31	1686	710	1.4	1.46	7.12
2800–3000	732	771	1	1.49	9.27	1068	735	1.33	1.49	8.17

Recent publications (Mason et al., 2017; Salem et al., 2019) have demonstrated that in convergent margins, sedimentary carbonates significantly contribute to the carbon budget of active arc volcanoes. About 75% of the emitted carbon would derive from crustal carbonate (Mason et al., 2017). The assimilation of carbon from crustal fluids and metasedimentary rocks has also been observed in continental large igneous provinces (Black and Gibson, 2019). The compilation of melt inclusions and gas emissions data from various tectonic settings highlights the importance of the presence of a crustal reservoir on the exogenic volatile flux budget (Fig. 5). Despite possible degassing effects during magma ascent, melt inclusions in komatiite – proxy for the primitive mantle (Sobolev et al., 2016), and melt inclusion in high-grade metamorphic rocks define two distinct endmembers (Fig. 5a). The primitive mantle, and to some extent mid-oceanic ridge basalts (MORBs), have a low water and low carbon content, unlike anatectic melts (Fig. 5a). The volatile content of plume, rift and arc volcanoes melt can then be interpreted as the results of mixing between the crust and the mantle. The same pattern is found when analysing the isotopic composition (He, C) of gas emission from active volcanoes and shear zones (Fig. 5b). Arc and rift settings display a mixed signature of both a mantle source and a low-metamorphic grade siliciclastic/carbonate source (e.g. Mason et al., 2017). Shear zone and faults might remobilise a significant amount of both high-grade crustal material and organic matter ($\delta^{13}\text{C} < -20$) during earthquakes. Thus, the presence of a metamorphic continental reservoir seems to have a long lasting effect on the global carbon cycle. Such fundamental question then links the evolution of Earth's fluid envelopes to the evolution of the crust, plate tectonics and the supercontinent cycle (Touret et al., 2016).

5.2. Implications for the evolution of plate tectonics

In this part, we discuss how the occurrence of nanorocks can help decipher the nature and evolution of geodynamic processes through time. The relative absence, up to date, of studies reporting nanorocks and COH(±N) fluid inclusions in terrains older than 0.6 Ga may be due either to (i) the preferential preservation of the continental crust, or (ii) a preservation issue (i.e. the inclusions originally present in the rock were then erased via as result of extensive rock re-equilibration and deformation) or (iii) a simple lack of data for this period. The first scenario is still disputed. Using the geochronological record, some studies have suggested that peaks in zircon abundance reflect an increase in continental growth rates (e.g. Condie, 1998; O'Neill et al., 2007). In contrast,

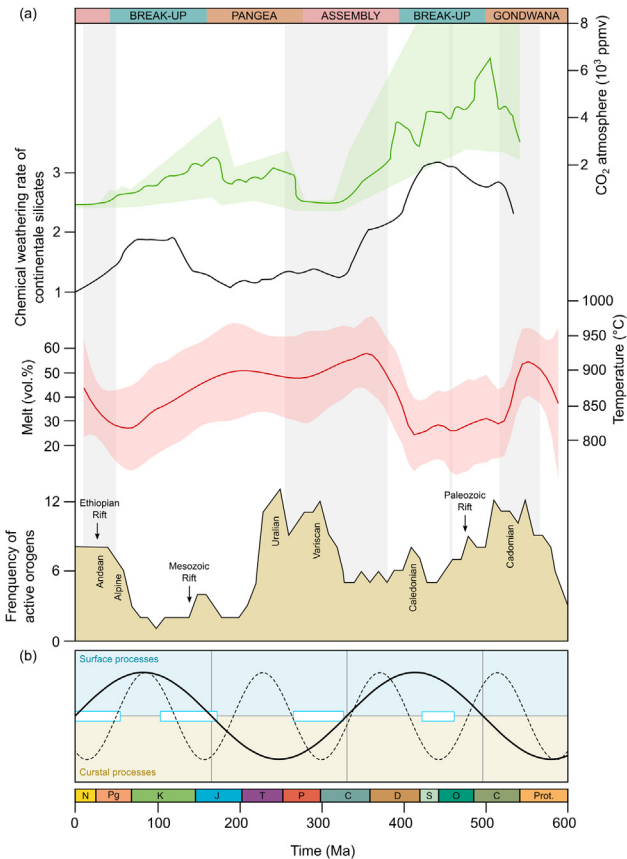


Fig. 4. Phanerozoic Wilson cycle showing the main trends in atmospheric and crustal evolution. (a) Atmospheric composition - GEOCARB III (Bernier and Kothavala, 2001). Chemical weathering rate of continental silicates, calculated using the exposed land area, the average surface temperature and the atmospheric CO₂ pressure. The curve is normalised to the present day (François et al., 1993). Average metamorphic temperature in the deep crust (>850 °C and 1 GPa - Fig. 1a, with 1σ error envelope) (Brown and Johnson, 2018) and melt production equivalent from Phanerozoic shales (vol.%). Number of active orogeny after Condie et al. (2015). Rift episodes after Hunt et al. (2017); Sánchez-García et al. (2019). Vertical shaded areas highlight the presence of primary MI and FI in the geological record. (b) Cyclic volatile reservoir transfers through the Phanerozoic (plain line); cosmic cycle (dashed line) and glacial epochs (rectangle) (Berger, 2013; Brink, 2019). Crustal storage alternate with surface storage over a period of ~320 Ma.

other studies have proposed that these peaks are contemporaneous with periods of supercontinent activity and are in fact an artifact related to the preferential preservation of certain magmas during collision cycles (e.g. Hawkesworth et al., 2009, 2010; Condie et al., 2011). However, a recent statistical treatment of the geochronological archive favours the episodic production of continental material, coupled with a proportional destruction (Puetz and Condie, 2020). The second is rather unlikely, as we find nanorocks in much older terranes with a complex metamorphic history. The third option is not as far-fetched as it may sound. Nanorocks were –and still often are– found in high grade terranes already extensively investigated for decades (e.g., Kerala Khondalite belt in India, Cesare et al., 2009; Ferrero et al., 2012) or even for centuries (e.g., Bohemian Massif, Ferrero et al., 2018b; Borghini et al., 2020). This suggests that they have been previously overlooked, possibly due to their small size, coupled with the unexpected oddity of having melt inclusions, typically a feature of igneous rocks, in metamorphic rocks.

Some studies proposed that self-sustaining subduction zones did not appear until ca. 2.0 Ga (Condie, 2018; Holder et al., 2019; Lenardic et al., 2019), others later in the Neoproterozoic (e.g. Stern, 2020). Until it did, the lack of tectonic feedback between the atmosphere and the solid Earth led to a net outgassing of water vapor, carbon dioxide and nitrogen into the primitive atmosphere and ocean (Shaw, 2008). While the loss of a significant part of the hydrogen to space led to the permanent oxidation of the Earth's surface (Catling et al., 2001; Zahnle et al., 2019), CO₂ and a part of H₂O present in the fluid envelopes were assimilated into the lithosphere through weathering processes (Bickle, 1996; West et al., 2002) and biological activity from 3.0 Ga onwards (i.e. oxygen photosynthesis – Lepot, 2020). The increase in carbon and hydrogen-saturated mineral species between 2.0 Ga and 3.0 Ga (Hazen et al., 2019) and the slow transition from stagnant-lid to mobile-lid tectonics can be then interpreted as the result of the increasing pairing between surface processes and deep Earth mechanisms (Valley et al., 2005; Sobolev and Brown, 2019).

Since 3.0 Ga critical changes in the thickness and thermal state of the crust, degree of crustal reworking and partial melting and the compositional variability of the crust (e.g. Condie, 1993; Veizer and Mackenzie, 2004; Nicoli et al., 2016; Brown and Johnson, 2018; Nicoli and Dyck, 2018; Palin et al., 2020), are thought to have led to a different, more efficient recycling method and long-term storage of volatile in the mantle through the generalisation of subduction zones (Newton, 1987; Santosh and Omori, 2008; Touret et al., 2016; Condie, 2018). Moreover, with the increase in continental volume and orogenic activity since 2.9–3.0 Ga (e.g. Dhuime et al., 2012; Condie et al., 2015), the volatile budget of the continental crust might have also grown with time (Santosh and Omori, 2008; Palya et al., 2011). Thus, the progressive storage of H, C and N in the solid Earth from 3.0 Ga onwards (e.g. Campbell and Taylor, 1983; Boerner et al., 1996; Berner and Berner, 2004; Touret et al., 2016) might have conjointly modified the physicochemical properties of Earth's solid and fluid envelopes (Korenaga, 2018). This could have launched the Earth system in a global feedback loop between deep Earth mechanisms and surface cycles (Goldblatt et al., 2009; Kerrich et al., 2006; Nisbet and Fowler, 2011).

The intrinsic link between the composition of the source (i.e. major and trace element chemistry and volatile content), the presence crustal derived magmas (e.g. Clemens et al., 2011; Nicoli et al., 2017; Villaros et al., 2009) (Fig. 2b, c) and graphite deposits (Luque et al., 2014) might suggest that some periods of Earth's history are more prone to the reworking of supracrustal lithologies than others (Campbell and Taylor, 1983). As it has been argued for the occurrence of graphite in the middle to lower crust (Oliver, 1992; Boerner et al., 1996; Wannamaker, 2000), the occurrence of pri-

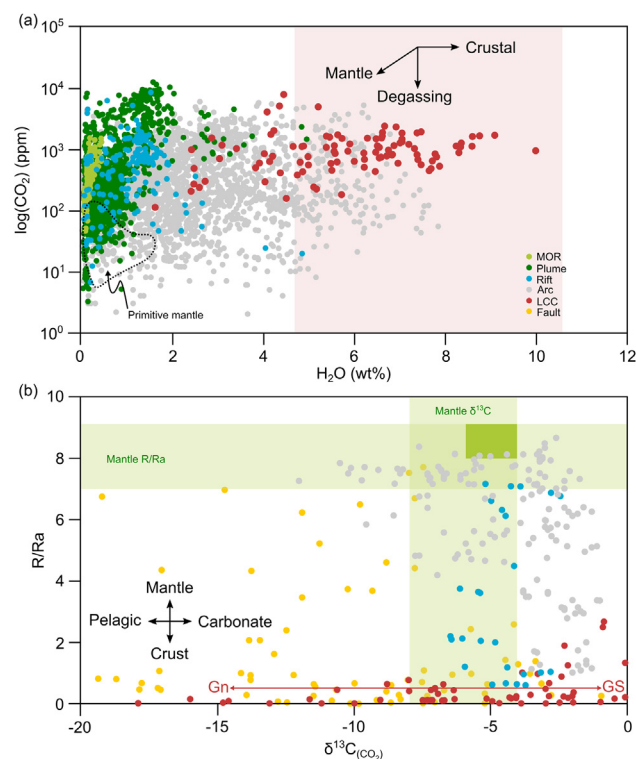


Fig. 5. Crustal input in volatile outgassing. (a) Measured volatile content (CO₂, H₂O) in melt inclusion in different geological settings (see Supplementary Data). MOR (mid-oceanic ridge) and nanorocks in the lower continental crust (LCC) represent two well-defined mantle and crustal endmembers respectively. The volatile content of plume (hot spot), arc volcanoes and continental rift might result from combination of mantle and crustal melt. The shaded area represents the range of H₂O value in nanorocks with no matching CO₂ values. Primitive mantle composition from komatiite melt inclusions. (b) Isotopic composition of volcanic and metamorphic gas (same colour code as Fig. 3a). R/Ra: ³He/⁴He in the sample (R), normalised to that in the atmosphere (RA = 1.38 × 10⁻⁶). δ¹³C reports carbon isotopes as the ratio of the heavy ¹³C isotope relative to the lighter ¹²C isotope relative to a standard in ‰. GS–Gn arrows indicates the range of δ¹³C composition found in metamorphic rocks from greenschists (GS) to Gn granulite (Gn) facies conditions. Fault represents the outgassing of the upper to middle crust caused by earthquake.

mary MI and FI in the geological record could directly evidence the efficiency of the exchange and redistribution of volatiles between the surface and the crust. As most of the MI and FI are younger than 850 Ma, one could link their occurrence to the presence of well-established plate tectonic mechanisms (Brown, 2006; Brown et al., 2020; Palin et al., 2020). It is worth noting that outside the Phanerozoic and late Neoproterozoic, MI and FI in metasedimentary rocks mainly occur between 2.5 Ga and 3.0 Ga (Fig. 2b and 3a), a period interpreted by some as the transition from a stagnant-lid regime to a mobile regime (Condie, 2018 and references therein). Additionally, and despite the lack of data on inclusions between 1.6 Ga and 2.1 Ga, the relatively high abundance of S-type granites (Fig. 2c) suggests that Paleoproterozoic terrains should contain a significant amount of nanorocks and primary COH(±N) fluid inclusions. This last period corresponds to the formation of the Columbia-Nuna supercontinent, characterised by an intense orogenic activity (Condie et al., 2015), and contains the first evidence of cold, long-lasting subduction zones (e.g. Weller and St-Onge, 2017; Holder et al., 2019; Wan et al., 2020). On the other end, the lack of data for the Boring Billion (0.85–1.7 Ga) could also be explained by the presence of single lid tectonics (Stern, 2020). Little continental reconfiguration and stable atmospheric and oceanic conditions (Roberts, 2013; Condie et al., 2015) would translate into limited transfer of volatiles from the surface to the

deep crust and the mantle, as indicated by the abundance of dry magmas during this period (Ashwal, 2010; Condie et al., 2014).

Therefore, the current distribution of MI and FI in metasedimentary rocks might argue for some degree of non-uniformitarianism in the evolution of plate tectonics. Further work is needed to complete the database and explore the different geodynamic scenarios mentioned above.

6. Conclusion and perspectives

The presence of preserved anatectic melt and volatile elements in inclusion in high-grade metamorphic mineral constitutes an incredible archive to investigate partial melting processes and crustal differentiation. The re-investigation of high-grade metamorphic rocks with the precise aim of finding preserved primary inclusions of deep fluids, coupled with the recent improvements in analytical techniques have showed that the amount of COH($\pm$ N) fluids in the lower crust might actually be higher than previously expected. However, while the volatile content of MI has been extensively studied in volcanic rocks, only a few studies focused on the origin and the implications of COH($\pm$ N)-fluids in MI and FI inclusions in high-grade metamorphic terranes for the evolution of geodynamic processes.

In the light of our findings, we argue that the lower crust might constitute an essential, long-term, volatile (i.e. H₂O, CO₂) storage unit, capable of influencing the composition of the surface envelopes through the mean of weathering, crustal thickening, partial melting and crustal assimilation during volcanic activity. Such hypothesis has implication beyond the scope of metamorphic petrology as it links geodynamic mechanisms to habitable surface conditions. Although most metamorphic studies focused on water and carbon dioxide, there is a still a large range of other volatile elements (e.g. S, N, Cl, F) that remain to be investigated in the context of preserved MI. Further constraints on the behaviour of volatile elements within the deep crust will be provided by future MI and FI studies in metamorphic rocks and will surely yield important new information on exchanges between reservoirs, giving greater insight into Earth's cycles.

In conclusion, that of the MI and FI in metamorphic rocks is a rich but still relatively uncharted realm. Its potential is vast but still partially untapped, as extensively discussed in the present work. In the near future, a concerted research effort should aim to find and characterise new instances of pristine inclusions in periods of the Earth's history currently underrepresented in our database, e.g. the Boring Billion. The merging of the messages of thousands of minuscule droplets of fluids trapped in the deepest roots of the continental plates will then eventually provide a truly comprehensive portrait of how the Earth's evolution proceeds through geological time.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.gsf.2021.101188>.

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