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PARTIAL MELTING AND P-T EVOLUTION OF ECLOGITE-FACIES METAPELITIC MIGMATITES FROM THE EGERE TERRANE (CENTRAL HOGGAR, SOUTH ALGERIA)

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15 ABSTRACT

16 The Egéré terrane (Central Hoggar, South Algeria) includes mafic eclogite lenses boudinaged in metapelitic rocks with high-pressure relicts. These metapelites show 17 18 textural records of partial melting, mainly primary melt inclusions enclosed in garnet 19 crystals and later crystallized as "nanogranitoids". Garnet porphyroblasts also contain 20 inclusions of quartz, kyanite, phengite, biotite, staurolite and rutile, and show a smoothed 21 prograde zoning with a Mn bell-shaped profile. The peak high-pressure metamorphic 22 assemblage consists of garnet, kyanite, phengite (Si up to 6.36), quartz, rutile, ±ilmenite, 23 ±feldspars and melt. Phengite has partially transformed into fine-grained aggregates of 24 biotite, plagioclase and K-feldspar, a microstructure interpreted as resulting from a 25 dehydration melting during exhumation. Phengite breakdown, along with other 26 retrograde reactions, produced a late paragenesis with biotite, plagioclase, K-feldspar, 27 quartz, almandine-rich garnet, ±sillimanite, ±staurolite, ±muscovite and ilmenite. The 28 thermodynamic modeling of *P*-*T* pseudosections allows to constrain various steps of the 29 metamorphic history: beginning of the garnet growth at 4.0 kbar and ~600°C during 30 prograde metamorphism; pressure peak at 14–20 kbar; temperature peak at 800–820°C; 31 formation of the last assemblage at 6.0–5.5 kbar and 725–685°C. Partial melting likely 32 started during the prograde path when crossing the H₂O-saturated solidus, at $T \ge 650$ -33 670°C and $P \ge 10$ kbar, continued upon heating, up to the peak conditions, as well as during decompression. This evolution is interpreted in terms of subduction of the 34 35 continental crust to mantle depths, followed by an exhumation through a clockwise P-T 36 path, during the Pan-African orogeny. The Egéré metapelites are relatively well-37 preserved eclogite-facies rocks, contain inclusions of "nanogranitoids" hitherto very little known in eclogite-facies metamorphic rocks, and represent an unusual trace ofsubduction within a Neoproterozoic orogen.

Keywords: Hoggar, Egéré, high-pressure metapelites, phengite, partial melting,
 nanogranitoids, pseudosections.

42 **1. INTRODUCTION**

The study of anatexis and *P-T* conditions of high-grade rocks is fundamental for understanding the geodynamic evolution and rheological behavior of the continental crust, particularly in convergent plate boundaries (*e.g.*, Rosenberg and Handy 2005). Understanding the significance of anatexis requires the coupling of detailed petrological, (micro)structural, geochemical and geochronological studies on migmatites.

This work combines two modern techniques to retrieve information on the petrogenesis of Neoproterozoic anatectic continental crust exposed in the Hoggar orogen, a Pan-African collisional belt extending from southern Algeria to northwestern Niger (Black et al. 1994; Caby 2003; Liégeois et al. 2003): (a) thermodynamic modeling of P-Tpseudosections and (b) microstructural study of melt-bearing inclusions, including nanogranitoids (*i.e.*, crystallized melt inclusions) and providing information on the relative timing and nature of anatexis.

55 While melt inclusions are well-known to igneous petrologists and have been 56 extensively studied in subvolcanic and extrusive rocks, their occurrence in high-grade 57 metamorphic rocks of anatectic terranes is a novel and powerful research tool (Cesare et 58 al. 2009, 2015). Melt inclusions in migmatites and granulites are generally trapped by 59 peritectic minerals produced by incongruent melting reactions and, hence, represent snapshots of primary anatectic melts (Cesare et al. 2009, 2015; Ferrero et al. 2012, 2018; 60 Bartoli et al. 2014, 2016). Although the study of melt inclusions in metamorphic rocks is 61 still in its infancy, those investigated so far indicate that the retrieved melt compositions 62 63 can be representative of the matrix melt present in the rock during entrapment (Acosta-64 Vigil et al. 2010, 2012; Cesare et al. 2015; Bartoli et al. 2016).

65 An increasing number of high-grade metamorphic rocks have been recently discovered and investigated from different localities in the Hoggar. Among them, rocks 66 67 that reached eclogite-facies conditions and preserved high-pressure (HP) parageneses are particularly interesting because their study is essential to unravel the nature and evolution 68 69 of orogenic processes (e.g., Berger et al. 2009, 2014; Doukkari et al. 2014, 2015; Adjerid 70 et al. 2015; Fettous 2016). To date, however, most of the reported and investigated HP 71 rocks are represented by metric to hectometric mafic eclogitic lenses. This study, in 72 contrast, investigates HP migmatitic metapelites of the Egéré-Aleksod terrane, Central 73 Hoggar, which are the first reported in the Hoggar with melt inclusions, now crystallized

into nanogranitoids. The principal goal of this contribution is elucidating the
metamorphic and anatectic history of these migmatites, and their implications for the
history of the Egéré terrane. This new data is combined with data from the surrounding
domains, to add new constraints on the eclogite-facies metamorphism in the Central
Hoggar and understand the geodynamic evolution of this orogenic belt.

79 2. GEOLOGICAL BACKGROUND AND FIELD RELATIONSHIPS

80 The Tuareg shield (S Algeria, W Mali and NW Niger) represents a Pan-African belt 81 located in between the West African Craton (WAC) to the west and the Saharan 82 metacraton to the east (Fig. 1). Formed by amalgamation of up to twenty-five terranes, it 83 is composed of Archean, Paleoproterozoic and juvenile Neoproterozoic rocks variably 84 reworked during the Pan-African orogeny (750-550 Ma; Black et al. 1994; Caby 2003; 85 Liégeois et al. 2003; Peucat et al. 2003; Liégeois et al. 2013; Liégeois 2019). According 86 to Liégeois et al. (1998), the long orogenic process includes a first major collision 87 between the Saharan metacraton and Eastern Hoggar terranes at ~750 Ma, marked by major thrusts and the exhumation of HP metamorphic rocks. At around 650 Ma, 88 89 subduction and the maximum of tonalite-trondhiemite-granodiorite (TTG) magmatism took place (Bechiri-Benmerzoug 2009). This episode produced HP metamorphism in the 90 91 Central Hoggar, the oblique collision of the Western Hoggar terranes with the WAC, and 92 northward migration of the Hoggar terranes along north-south shear zones, hundreds of 93 kilometers long (Bertrand and Caby 1978; Bertrand et al. 1986; Black et al. 1979; Caby 94 et al. 1989; Peucat et al. 2003). Later events, between 630 and 580 Ma, included high-95 temperature (HT) regional metamorphism and the generation of high-K calc-alkaline 96 batholiths (Liégeois et al. 2003; Acef et al. 2003; Abdallah et al. 2007).

97 The polymetamorphic Central Hoggar is well-exposed at the core of the Tuareg 98 shield (Fig. 1a). It is made of eight terranes, delimited by shear zones: Laouni, Aouilène, 99 Tefedest, Egéré-Aleksod, Azrou n'Fad, Assode-Issalane, Serouenout and Tazat (Black et 100 al. 1994). The first five are grouped under the "LATEA metacraton" acronym (Fig. 1b; 101 Liégeois et al. 2003, Liégeois 2019).

102 Lelubre (1952) and Duplan (1967, 1972) were the first to study the Egéré-Aleksod 103 terrane, where Latouche (1985) described ortho- and para-derived rocks constituting 104 domes and basins, respectively. However, Doukkari et al. (2014, 2015), Doukkari (2016), Arab et al. (2015) and Arab (2016) have considered the Egéré terrane as composed of two 105 106 juxtaposed tectonic units separated by a mylonitic thrust, considered as a major tectonic contact: (i) The Arechchoum basement made of migmatitic ortho- and para-gneisses 107 108 derived from Paleoproterozoic protoliths; (ii) the Egéré meta-sedimentary cover, an 109 eclogite-facies unit made up mainly of kyanite-bearing metapelites (Fig. 2c-d) associated 110 with subordinate marbles, quartzites, several-meter-long lenses of mafic eclogite, amphibolite and rare ultramafic rocks (Fig. 2a-b; Sautter 1985; Doukkari et al. 2014, 2015; Arab et al. 2015). In the investigated Izzilatène area, migmatites appear as metatexites (Fig. 2d), and mafic eclogites constitute lenses within both migmatites and marbles (Fig. 2a-b). Field observations, metamorphic constraints and U-Pb ages, as well as geochemical data, suggest that the Egéré represents a former passive continental margin intruded by mafic rocks, which afterwards was deeply subducted and then exhumed (Arab 2016; Doukkari 2016; Ouzegane et al. in preparation).

118 **3. PETROGRAPHY AND MINERAL CHEMISTRY**

119 Representative metapelitic samples were collected from the Egéré meta-sedimentary 120 unit in the Izzilatène area, at the north of the Egéré terrane and ca. 50 km south of similar 121 rocks first reported by Arab et al. (2015) in the Tighsi area (Fig. 1b). In the field, these 122 metapelites appear as stromatic migmatites, with cm-to-mm-thick leucocratic bands 123 (leucosomes) alternating with darker bands rich in garnet and biotite, and parallel to the 124 main foliation. In hand specimen and under the microscope, they are banded rocks, 125 consisting of two petrographically different domains, leucosomes and melanosomes. The 126 leucosomes consist of quartz, feldspars and minor garnet. They are interspersed with 127 quartz-poor mafic melanosomes, which contain large garnet crystals, up to several 128 centimeters in diameter, together with kyanite, biotite, white micas, staurolite, sillimanite, 129 rutile and ilmenite (Fig. 2c-d).

130 Eight specimens of migmatitic metapelites (Table A in supplementary materials 131 [SM]) have been studied under the SEM and analyzed at the electron microprobe (see 132 methods in SM). They consist of variable proportions of garnet, kyanite, white micas 133 (phengite and muscovite), biotite, K-feldspar, plagioclase, guartz, staurolite and 134 sillimanite. Under the polarizing microscope, these rocks show a fine- to medium-grained matrix made of quartz, feldspars, phengite, biotite, kyanite, sillimanite and staurolite, 135 136 which encloses cm-sized porphyroblasts of garnet and, less commonly, of kyanite (Fig. A 137 in SM). Accessory minerals are apatite, rutile, allanite, tourmaline, ilmenite and zircon. 138 These rocks often show an *augen*-like microstructure produced by large monocrystalline 139 quartz crystals or polycrystalline quartz aggregates (Fig. 3e), and/or large polymineralic 140 pseudomorphs after phengite (Figs. 3f, 4d-f; see below).

All the rocks sampled show comparable features; on the basis of microstructural
observations, the minerals have been grouped into the following three main metamorphic
parageneses.

144(i) The earliest preserved assemblage (M_1) is represented by abundant mineral relicts145armored within garnet porphyroblasts (Grt_1) , including quartz (Qz_1) , biotite146 (Bt_1) , phengite (Ph_1) , staurolite (St_1) , kyanite (Ky_1) , plagioclase (Pl_1) , rutile147 (Rt_1) and ilmenite (IIm_1) (Figs. 3a-d, 4c).

- (ii) The high-grade peak paragenesis (M₂) consists of the rims of the garnet
 porphyroblasts (Grt₂) at contact with their matrix, made of kyanite (Ky₂),
 phengite flakes (Ph₂), quartz (Qz₂), plagioclase (Pl₂), K-feldspar (Kfs₂),
 apatite, rutile (Rt₂) and ilmenite (IIm₂) (Fig. 4a). The minerals in the matrix,
 mainly Qz₂, Ph₂ and Ky₂, define the main foliation in the rock. The
 coexistence of melt with this paragenesis is discussed later.
- (iii) A late assemblage (M₃) occurs either as a recrystallized matrix with biotite (Bt₃), plagioclase (Pl₃), K-feldspar (Kfs₃), quartz (Qz₃), muscovite (Ms₃), garnet (Gt₃), sillimanite (Sil₃), ilmenite (Ilm₃) and/or staurolite (St₃) (Figs. 3g-h, 4d), or as fine-grained symplectites made up of Bt₃, Pl₃ and Kfs₃ (Fig. 4e-f), which formed mostly after Grt₁₋₂ and Ph₂.
- 159 In the following subsections, we describe the microstructure and composition of the 160 main minerals, whose analyzes are summarized in Tables B-F (SM).
- 161 *3.1. Garnet*

162 Three generations of garnet have been distinguished on the basis of microstructures 163 and compositions. The first two generations $(Grt_{1,2})$ correspond to the core and rims, 164 respectively, of euhedral to subhedral porphyroblasts, ~ 0.4 to 7 cm in diameter. These porphyroblasts are well preserved, except some rims that appear partially replaced by 165 166 fine-grained Bt₃+Pl₃ aggregates (Figs. 3a, 4b). These Grt₁₋₂ crystals are crowded with 167 mono- and poly-crystalline inclusions, which often define an either straight (Fig. Ac, e, g 168 in SM), sigmoid or spiral (Fig. Ab) internal foliation, at variable angles with respect to the foliation in the matrix but occasionally in continuity with it, indicating that the garnet 169 170 porphyroblasts grew before or during the main deformation event. Some inclusions are 171 large (0.5–1.5 mm), anhedral, and consist of either individual or polycrystalline 172 inclusions of Qz₁, Ph₁, Bt₁, St₁, Rt₁, Ilm₁ and/or zircon (Fig. 3b-d). Others, from 0.1 to 0.5 173 mm in size, are characterized by the presence of a thin optically-continuous moat of 174 plagioclase surrounding quartz (Pl in Figs. 3c, 4c). There are also small (~5-30 um) 175 polymineral inclusions with the negative shapes of the host garnet (Fig. 5), which have all 176 the characteristics of nanogranitoids (Cesare et al. 2015). The largest polymineral 177 inclusions are visible mainly in the mantle and rim of the garnet porphyroblasts, while the 178 nanogranitoids are apparently distributed throughout the entire crystals. Sometimes, 179 garnet is also skeletal and appears intergrown with quartz (Fig. 3e).

Garnet porphyroblasts are rich in almandine (Alm₈₃₋₆₅Prp₄₋₁₃Grs₂₆₋₈Sps₆₋₀: Fig. 6a-d; Table B in SM). In most of the samples, they show profiles characterized, from core to rims, by decreases in X_{Grs} , X_{Sps} and X_{Fe} and an increase in X_{Prp} (Fig. 6a-d). A late compositional zoning, with increases in X_{Fe} , X_{Sps} and X_{Grs} , and a decrease in X_{Prp} (Fig. 6c), is observed in the outer rims that are corroded by a late Bt₃+Pl₃ symplectite.

Grt₃ is observed as small crystals either scattered in the matrix, in association with Bt₃, Pl₃, St₃ and sometimes Sil₃, or close to the large Grt₁₋₂ porphyroblasts, of which they can represent modified relics of their resorbed rims. Compared with Grt₁, Grt₃ is richer in Fe and poorer in Ca (Table B), having a very low grossular content ($X_{Grs} = 0.06-0.12$) and a fairly high X_{Fe} (0.88–0.94), similarly to the retrograde outer rims of some corroded garnet porphyroblasts (Fig. 6c-d).

3.2. White micas

192 Three generations of white mica have been observed. Phengite Ph_1 appears as rare 193 inclusions in the garnet porphyroblasts. Phengite Ph₂, by far the most abundant, occurs in the matrix as large flakes, $\sim 0.5-3.0$ mm long, defining the main foliation; it therefore 194 195 developed before and/or during the main deformation event. These large crystals 196 frequently show Bt_3 and Ilm_3 lamellae, exsolved preferentially parallel to the (001) 197 cleavage (Figs. 3f, 4e-f). Ilm₃ exsolution lamellae are particularly abundant in the vicinity of pre-existing Ilm₂ crystals (1 in Fig. 4f), suggesting that the initial Ph₂ was more 198 199 titaniferous here. Ph₂ flakes are partially or totally replaced by fine-grained symplectitic 200 aggregates of Bt₃+Pl₃, bounded by a continuous Kfs₃ corona at contact with Oz₂ (Fig. 4f) and by a Pl₃+Ilm₃ symplectite around Ilm₂ (2 in Fig. 4f). Similar pseudomorphs after 201 phengite, observed elsewhere, are generally interpreted as formed during exhumation 202 203 (e.g., Indares and Dunning 2001; Korhonen and Stout 2005; Lang and Gilotti 2007; 204 Dokukina et al. 2017). The third generation of white mica consists of small grains (0.1– 0.3 mm) of muscovite Ms₃, some of which are by-products of the Ph₂ breakdown, being 205 isolated in the Bt₃+Pl₃ symplectites (Figs. 3f, 4f). 206

207 White micas in most microstructural settings have a phengitic composition, though 208 with various Si contents (Fig. 4e; Table C in SM). Ph₁ is poorly substituted (Si ≈ 6.25 209 a.p.f.u.). The Ph₂ flakes are zoned, showing a decrease in phengite substitution from core to rims: Si = 6.44 a.p.f.u. at core vs. 6.05 at rim; $Al^{t} = 4.96$ vs. 5.76; Ti = 0.17 vs. 0.08; 210 (Mg + Fe) = 0.51 vs. 0.22. The initial composition of a Ph₂ zone rich in Ilm₃ exsolution 211 212 lamellae (1 in Fig. 4f) was reconstructed at the electron microprobe by scanning ~100um² adjacent areas with the electron beam during acquisition; the average composition 213 (Ti-Ph in Table C) suggests that Ph₂ before exsolution was here strongly substituted by 214 215 the theoretical end-member Ti-Fe-phengite ($\sim 1/3$ of K₂Fe₂Ti₂[Si₆Al₂]O₂₀(OH)₄: 216 Auzanneau et al. 2010). Ms₃ in the Bt₃-Pl₃ pseudomorphs after phengite Ph₂ is similar in 217 chemistry to individual Ms_3 grains present in the rock matrix (Table C; Fig. 4e; Si = 218 5.96-6.05 a.p.f.u.; Ti = 0.03-0.12; [Mg + Fe] = 0.16-0.20) and to the rims of Ph₂ flakes, 219 suggesting that the zoning of the latter is linked to the formation of the Bt₃-Pl₃ 220 symplectite and thus to retrogression.

3.3. Biotite

222 Biotite occurs in four main microstructural varieties: (i) As inclusions in garnet (Bt₁: 223 Fig. 3b). (ii) In the rock matrix, associated with Pl₃, Kfs₃ or St₃. Commonly, these biotite-224 rich aggregates are fairly coarse-grained and oriented according to the main foliation 225 (Figs. Ac in SM, 3g bottom), but they sometimes show a gradual transition with fine-226 grained unoriented Bt₃+Kfs₃+Pl₃ aggregates after Ph₂ (Fig. 3g upwards). Therefore, 227 biotite in the matrix is a post-peak Bt₃, the size and orientation of which depend on when 228 it formed relative to the main deformation event. (iii) Bt_3 is also intergrown Sil₃ (Fig. 3h). 229 (iv) Finally, minute Bt₃ lamellae parallel to the (001) plane of Ph₂ resulted from late 230 exsolution (Fig. 4e-f).

The large oriented biotite flakes in the matrix are unzoned, more magnesian ($X_{Fe} = 0.40-0.54$) and lower in Ti (≤ 0.34 a.p.f.u.) with respect to biotite from the fine-grained aggregates in the matrix (Bt₃: $X_{Fe} = 0.67-0.87$; Ti = 0.36-0.48 a.p.f.u.) (Fig. 6g; Table D in SM). Biotite exsolution lamellas in phengite are similar to biotite in the pseudomorphs after Ph₂ (Bt₃; $X_{Fe} = 0.66-0.69$; Ti = 0.32-0.34 a.p.f.u.) (Fig. 4f; Table D).

236 *3.4. Feldspars*

The rare plagioclase crystals (0.1 mm) present in the leucosomes are relatively homogeneous, with X_{An} ranging from 0.29 to 0.33 (Fig. 6h; Table E). The thin plagioclase coronae surrounding Qz₁ inclusions in garnet are oligoclase (An_{14–15}; Fig. 3c; Table E). The plagioclase Pl₃ associated with Bt₃, St₃ and/or Sil₃ in the matrix, and produced essentially from Ph₂ (Fig. 4e-f), is An_{14–26} (Table E), whereas Pl₃ in the Bt₃-rich aggregates that replaced garnet porphyroblasts (Figs. 3a, 4b) is slightly richer in anorthite (An_{25–41}), due to the Ca content of garnet from which it grew.

248 *3.5. Other minerals*

Kyanite is observed as inclusions in garnet (Ky₁) and as large laths in the matrix (Ky₂), ranging in size from ~0.1 mm to 2 cm, which, together with micas and quartz, defines the main foliation (Fig. Ag in SM). Ky₂ may appear partially or completely transformed to mm-sized bundles of fibrolitic sillimanite (Sil₃). Sil₃ appears also intimately intergrown with Bt₃ as sheaf-like clusters (Fig. 3h).

Staurolite occurs both as inclusions in Grt₁ porphyroblasts (St₁ in Fig. 3c-d; X_{Mg} = 0.13–0.15; Si = 7.4–7.6 a.p.f.u.; TiO₂ = 0.64–0.87 wt%) and in the matrix associated with Bt₃, Kfs₃, Pl₃, Gt₃ and Sil₃, as small scattered subhedral crystals (St₃ in Fig. 3g-h; X_{Mg} = 0.06–0.10; Si = 6.65–7.37 a.p.f.u.; TiO₂ = 0.49–0.74 wt%), with a more intense yellow

pleochroism. Clear Si₋₁Al^{IV}₊₁Al^{VI}₊₁Fe₋₁ (*i.e.*, Tschermak) and Mg₊₁Fe₋₁ substitutions are observed in the analyzed staurolites (Fig. 6f; Table F in SM).

Rutile largely replaced by ilmenite, fluorapatite, allanite, zircon and tourmaline are accessory minerals.

262 4. MICROSTRUCTURAL EVIDENCE OF PARTIAL MELTING

There are typical microstructures indicating the former presence of melt during highgrade metamorphism, provided that crystallized and/or solidified melt has not been affected by subsequent subsolidus deformation and recrystallization phenomena. The most obvious are those present in shallow contact metamorphic aureoles, where melt has either solidified into glass or crystallized into a granophyric intergrowth. Former melt commonly constitutes thin films in between most crystals, and may include anhedral reactant and/or euhedral peritectic minerals (Holness and Sawyer 2008).

270 The presence of melt is not so obvious in slowly cooled contact and regional 271 anatectic rocks (Sawyer 2001). There is a consensus however that several microstructures 272 are diagnostic: (i) subhedral to euhedral shapes, particularly for minerals that do not 273 typically produce crystal faces in metamorphic rocks (Vernon and Collins 1988); (ii) melt 274 pseudomorphs, identified by quartz or K-feldspar or sodic plagioclase, forming either 275 strongly cuspate grains or thin films between other crystals (Harte et al. 1991; Clemens 276 and Holness 2000; Sawyer 2001; Marchildon and Brown 2002; Holness and Sawyer 277 2008); (iii) structures indicative of melt-producing reactions, like euhedral peritectic 278 minerals, anhedral relicts of reactant minerals surrounded by melt pseudomorphs, 279 oscillatory zoning, symplectitic coronae of Bt + Pl or Bt + Qz around ferromagnesian 280 crystals (Sawyer 1999; Vernon and Collins 2011); (iv) magmatic flow orientation 281 underlined by undeformed euhedral elongated crystals (Paterson et al. 1989; Sawyer 282 2001; Vernon and Collins 2011).

283 In the case of the studied Egéré metapelites, the first indications of the presence of melt are based on field observations of leucosomes parallel to the main foliation. Under 284 285 the microscope, the leucosomes appear made of quartz with subhedral to euhedral crystals of sodic plagioclase and K-feldspar. The presence of several types of mineral 286 287 intergrowths also points to the presence of melt: (i) Bt₃+Pl₃ aggregates around garnet 288 (Figs. 3a, 4b), interpreted by some authors as resulting from the breakdown of garnet during cooling and melt crystallization (Janák et al. 1999); (ii) skeletal garnet intergrown 289 290 with quartz (Fig. 3e), interpreted as related either to prograde dehydration during melting 291 reactions or to a back-reaction with melt (Waters and Whales 1984; Waters 1988, 2001; 292 Vernon and Collins 1988; Powell and Downes 1990); (iii) Sil₃+Bt₃ sheaf-like 293 intergrowths (Fig. 3h), interpreted as products of a melt-residuum back-reaction (Dusel 294 and Foster 1983).

295 Strong microstructural evidence of the former presence of melt in high-grade 296 metamorphic rocks is also provided by some polycrystalline inclusions in garnet (e.g., Hartel et al. 1990; Perchuk et al. 2005; Lang and Gilotti 2007; Cesare et al. 2009, 2015). 297 298 Cesare et al. (2009, 2015) described what they termed "nanogranitoids", referring to tiny 299 polycrystalline inclusions, up to a few tens of micrometers in size, representing 300 crystallized droplets of primary anatectic melt, trapped by the host peritectic mineral 301 growing during an incongruent melting reaction. Lang and Gilotti (2007) described 302 relatively large and mostly anhedral inclusions in garnet, consisting of mono- or poly-303 crystalline quartz rimmed by a discontinuous moat of plagioclase and K-feldspar. They 304 interpreted this microstructure as representing a crystallized melt from the matrix, which 305 infiltrated or invaded the garnet crystals along cracks or grain boundaries, probably during the expansion due to the transformation into quartz of former coesite inclusions 306 307 (see also the crown-like wavy microstructure of Korsakov and Hermann, 2006). On the 308 basis of experiments on inclusion-bearing garnet crystals from mafic eclogites, Perchuk 309 et al. (2005) showed that some large polycrystalline inclusions with melt-bearing 310 microstructures could be produced by the post-entrapment melting of former solid 311 polycrystalline inclusions involving hydrous minerals.

312 In the specific case of the Egéré migmatites, all three types of polycrystalline 313 inclusions are present in garnet and demonstrate anatexis of the host rock:

- 314 (i) Nanogranitoids: These inclusions are very small (~5-30 µm), equant, and 315 show the negative crystal forms of the host garnet (Fig. 5). They are rare and scattered throughout the garnet porphyroblasts. These inclusions are 316 317 composed of quartz, biotite, plagioclase, K-feldspar, chlorite and accessory 318 minerals such as zircon and allanite. Given their distribution, size, shape and 319 mineralogy, they are interpreted as nanogranitoids. While zircon and allanite 320 would constitute solid inclusions entrapped with the melt, quartz, feldspars 321 and biotite probably represent daughter minerals crystallized from the trapped 322 melt; chlorite may be a low-T mineral formed during the retrograde P-T path.
- 323 (ii) Inclusions of quartz surrounded by a thin feldspar moat: Very similar to 324 those described by Lang and Gilotti (2007), these inclusions are subhedral to 325 euhedral, $\sim 0.1-0.5$ mm in size (Figs. 3c, 4c), and made up of mono- or poly-326 crystalline quartz (Qz_1) in the center, rimmed by a thin moat of plagioclase, 327 either optically continuous or sometimes polycrystalline (Fig. 4c), which 328 suggest the former presence of melt (Sawyer 2001; Lang and Gilotti 2007). 329 Although these inclusions have been observed mainly at the rims of garnet 330 porphyroblasts, the secondary infiltration of matrix melt might not be the only 331 explanation for this particular microstructure; a primary mixture of reactants 332 and melt-crystallization products could be an alternative interpretation.

Large polyphase inclusions: About 0.5 to 1.5 mm in size, they consist of
quartz, feldspars, phengite, biotite, kyanite, staurolite, rutile and ilmenite, and
are found in the intermediate and rim regions of garnet porphyroblasts. They
could be interpreted as phengite-bearing polycrystalline inclusions that
partially melted after their entrapment in garnet (Perchuk et al. 2005). A more
detailed and specific study would however be necessary to confirm such an
origin.

340 5. PHASE EQUILIBRIA MODELING

341 In order to specify the metamorphic evolution of the metapelites, *P*-*T* pseudosections 342 (Fig. 7) were calculated using the software THERMOCALC (Powell and Holland 1988) 343 and an updated release of the thermodynamic dataset of Holland and Powell (2011) (see 344 method in SM for the details). To model the early prograde *P*-*T* evolution, we have 345 constructed a pseudosection for sample T78 (Fig. 7a). Results should however be 346 considered with caution, for two main reasons: (i) The early prograde evolution of a 347 pelitic metasediment takes place reasonably under conditions of H₂O saturation. As soon 348 as partial melting occurs, H₂O is strongly partitioned into the melt and the aqueous fluid 349 rapidly becomes absent in the system. The quantity of H₂O that has been considered for 350 modeling this early stage, higher than the loss-on-ignition value of the final rock, takes 351 these considerations into account, but is nevertheless arbitrary which causes large 352 uncertainties on the high-T part of Figure 7a. (ii) Some melt may have been lost from the 353 analyzed volume of the rock, and in part segregated to form the leucosomes (e.g., White 354 et al. 2004; Lasalle and Indares 2014). It would be wise to reintegrate a certain (arbitrary) 355 amount of melt into the modeled composition. We actually preferred considering a 356 sample large enough to include a representative amount of leucosome.

357 During partial melting, any water initially present dissolves into the melt (see above). 358 This is why it is better to calculate H₂O-undersaturated pseudosections to model the peak 359 and retrograde P-T evolution of these rocks, which we did using the bulk composition of 360 samples T52 and T78, of which the H₂O content was deduced from their loss-on-ignition 361 value (Fig. 7b-c; see Table A in SM for the compositions). A melanocratic part of sample T52 was also separated and analyzed, in order to model a rock microdomain (Fig. 7d) 362 363 that mainly consists of Bt₃, St₃, Sil₃, Pl₃, Grt₃, quartz and ilmenite, which is considered to 364 be the most retrograde paragenesis.

365 6. DISCUSSION

366 On the basis of microstructural and mineralogical evidence, at least three distinct 367 stages of metamorphism have been recognized in the Egéré metapelites: (i) the earliest 368 metamorphic stage (M_1) is prograde and documented by mineral inclusions enclosed in

369 garnet porphyroblasts; (ii) the peak eclogite-facies stage (M_2) developed a paragenesis 370 made of garnet, kyanite, phengite, quartz, rutile and melt; (iii) finally, the retrograde post-371 peak stage (M_3) is highlighted by post-foliation minerals that developed in the matrix, 372 mainly as fine-grained pseudomorphs resulting from the breakdown of peak minerals like 373 phengite and garnet.

374 *6.1 Prograde evolution*

The prograde metamorphic evolution is documented by mineral inclusions preserved in the core of garnet porphyroblasts (Grt₁), including Qz_1 , Ky_1 , Bt_1 , St_1 , Ph_1 , Rt_1 and IIm_1 (Figs. 3a-d, 4c). Some of these inclusions exist alongside polycrystalline inclusions with negative crystal shapes that are interpreted as nanogranitoids (Fig. 5), and thus testify to the presence of a melt trapped in the host garnet during its growth (Section 4).

380 It is reasonable to assume that any initial prograde zoning in garnet would have been 381 smoothed or erased by diffusion at peak high-grade conditions (Spear and Florence 1992; 382 Caddick et al. 2010), which is corroborated here by the generally flat zoning pattern 383 observed in the core of the studied garnets (Fig. 6a, c). However, it is also well known 384 that bell-shaped X_{Sps} zoning profiles in garnet (e.g., samples T78, T138; Fig. 6a, c) 385 indicate prograde growth zoning (e.g., Tracy 1982; Loomis 1983; Mahar et al. 1997). 386 Likewise, the grossular content, higher in the core of garnet crystals than in their rims, is 387 also typical of a prograde growth zoning of garnet from metapelites (e.g., Woodsworth 388 1977; Chernoff and Carlson 1997; Keller et al. 2006). Overall, all these observations 389 suggest that Grt₁ grew at the expense of minerals now preserved as single mineral 390 inclusions, via reactions that occurred during the prograde evolution of the rock. At least 391 part of Grt₁ crystallized in the presence of melt, as shown by the presence of 392 nanogranitoids.

393 The prograde *P*-*T* trajectory was modeled by calculating a pseudosection for sample 394 T78 (Fig. 7), despite the uncertainties linked to such modeling (Section 5). Given the 395 high temperature reached by these rocks (800–820°C, see below), the initial prograde 396 zoning of garnet has been likely smoothed by diffusion at peak conditions. Hence, we did 397 not use garnet isopleths for modeling the prograde *P*-*T* path, which thus remains poorly 398 constrained. However, we tried to define a *P*-*T* evolution (red dashed arrow in Fig. 7a) 399 that explains both the growth of the garnet porphyroblasts (by crossing the garnet 400 isomodes: red dotted lines) and the entrapment of the various inclusions. An anchor 401 point, at ~600-630°C and ~6.0-6.5 kbar (blue ellipse in Fig. 7a), is provided by the 402 assemblage $Qz_1 + Bt_1 + Ph_1 + St_1 \pm Ky_1 + Pl + Ilm_1 (+Grt_1)$, shielded as inclusions in Grt_1 403 porphyroblasts of sample T78; it is further constrained by the intersection of Bt₁ and St₁ isopleths (x(Bt) = 0.62-0.64; x(St) = 0.85-0.88). To account for the presence of 404 405 "nanogranitoids" in garnet porphyroblasts, this early P-T evolution must have reached 406 fields of the pseudosection where melt ("Liq") is present and where the isomodes of 407 garnet are tightly spaced suggesting its peritectic growth (Fig. 7a). This could have 408 occurred during the crossing of the H₂O-saturated rock solidus at ca. $T \ge 650-670^{\circ}$ C and 409 $P \ge 10$ kbar. A modeled melt formed along the postulated trajectory at 10 kbar and 410 670°C, has the following composition: Si_{39.84}Al_{10.15}Fe_{0.12}Mg_{0.06}Ca_{0.38}Na_{7.38}K_{1.69}O₁₀₀. 411 xH₂O, which, after crystallization, would give a mineral assemblage compatible with the 412 nanogranitoids observed (Fig. 5; Section 4): 62.8 Pl (Ab₉₅) + 22.0 Qz + 13.8 Or + 1.4 Bt 413 (in vol.%; molar volumes from the Thermocalc database).

414 *6.2 Peak P-T conditions*

The prograde evolution should have continued towards the peak of metamorphism. The H₂O-undersaturated pseudosections (Section 5) are used to constrain the peak or near-peak *P-T* conditions (Fig. 7b-c). The peak assemblage (M₂) observed in the matrix (Grt₂ + Ph₂ + Ky₂ + Qz₂ + Rt₂ + Ilm₂ + Kfs₂ + Pl₂ ± Liq) and whose feldspars are concentrated in leucosomes, is stable over a large *P* range between 12 and 20 kbar. It is therefore necessary to better define the peak conditions by using the isopleths of garnet and/or phengite. In doing so, several drawbacks arise:

422 (i) The temperatures reached during the peak of metamorphism have produced a 423 significant, but incomplete, chemical homogenization of garnet porphyroblasts, with 424 almost flat zonation profiles except at their rim, which reequilibrated during the 425 retrograde metamorphism (Fig. 6a-d). Hence, it is appropriate to use the composition of 426 the large central parts of the garnet porphyroblasts to refine the peak P-T conditions. 427 However, the calculated X_{Fe} and X_{Grs} isopleths in garnet (x(Grt) and z(Grt) in Fig. 7b-c) 428 vary only very little over the large P range in which the paragenesis M₂ is stable, and are 429 thus of limited interest.

(ii) The composition of phengite seems more reliable because the isopleths *Si(Ph)*, related to the content in celadonite, are closely spaced (Fig. 7c-d), and can be applied to the nearly homogeneous Ph₂ flakes. The measured values of Si in phengite suggest P >14 kbar when compared with the calculated *Si(Ph)* isopleths (*Si(Ph)* ± 1 σ : 6.278 ± 0.038 [T78]; 6.234 ± 0.022 [T52]). High-grade conditions are also indicated by the Ti richness of parts of Ph₂ (Section 3.2; Ti-Ph in Table C; *e.g.*, Nahodilová et al. 2014; Auzanneau et al. 2010).

437 (iii) The liquid produced by anatexis (Section 4) could have been partially extracted 438 from the rock, producing a change in composition of the system and a shift of the solidus 439 curve towards higher temperatures (dashed blue arrows in Fig. 7b-c). In the borderline 440 case where all melt is extracted, the solidus curve would shift and follow the P-T441 conditions of the rock, since the solidus corresponds to those conditions where the melt 442 exists only in infinitesimal amount. It is unlikely, however, that all of the melt was 443 extracted because a critical threshold of liquid is required for melt interconnection and

migration (<1 to a few vol.%; Laporte and Watson 1995; Laporte et al. 1997; Rosenberg and Handy 2005). Consequently, the modeled pseudosections of Figure 8b-c give a "snapshot" that probably approaches but does not reach this borderline case. They would reflect the situation close to the T peak, and are thus poorly suited for determining the conditions prior to it.

449 The *P* peak must have been reached at lower temperatures relative to the ultimate 450 solidus curve given by the pseudosections, in the stability field of the M₂ paragenesis (Grt + Ph + Ky + Oz + Ilm \pm Rt + Kfs + Pl), and in the wide range of P between 14 and 20 451 452 kbar given with great uncertainty by the garnet and phengite isopleths (Fig. 7b-c). Some 453 constraints, like Ph₂ still abundant in the rock, together with feldspars and melt, and the 454 location of the ultimate solidus curve, suggest a subsequent T peak at about 800–820°C. 455 The Ti-in-Zrn thermometry applied to similar neighboring metapelites yielded peak temperatures of $811 \pm 15^{\circ}$ C (Bruguier et al. 2014). The *P-T* trajectories drawn in Figure 456 457 8b-c are an attempt to represent this imperfectly constrained evolution.

458 *6.3 Retrograde evolution*

The *P-T* trajectory of exhumation is documented by the M_3 paragenesis, made of abundant Bt₃, Pl₃, Kfs₃ and quartz, scarce Sil₃, Ms₃, Grt₃, ±St₃ and Ilm₃. The M₃ assemblage often forms unoriented fine-grained polymineralic aggregates (almost symplectites) and therefore postdates the main foliation. In some specimens, however, Bt₃ shows a significant orientation (Section 3.3; Fig. 3g), indicating some deformation at this stage. M₃ developed at the expense of previous M₂ minerals, mainly garnet and phengite. Sillimanite and ilmenite also replaced kyanite and rutile, respectively.

466 The garnet porphyroblasts, largely homogenized at the peak of metamorphism (see 467 above), show a drop in X_{Fe} and X_{Grs} and an increase of X_{prp} at their rims ("rims" in Fig. 468 6a, c). Moreover, when they have been corroded by a fine-grained Bt₃+Pl₃ aggregate 469 (Fig. 4b), these porphyroblasts show slight increases of X_{Fe} and X_{Grs} at their very edges ("outer rim" of Fig. 6c). Some small anhedral Grt₃ crystals, whose X_{Fe} is even greater 470 471 (Fig. 6d; Table B), may well be relicts of these porphyroblast edges, isolated during the 472 development of the replacing minerals. Many authors have interpreted such an increase in 473 X_{Fe} as resulting from Fe-Mg exchange between garnet and surrounding biotite during 474 retrogression (e.g., Tracy et al. 1976; Woodsworth 1977; Tuccillo et al. 1990; Konrad-475 Schmolke et al. 2005).

The phengite flakes in the matrix (Ph₂) have been partly or totally replaced by a finegrained Bt₃+Pl₃ aggregate, which is further rimmed by a Kfs₃ corona at contacts with Qz₂ (Figs. 3f, 4e-f). They show a chemical evolution towards the muscovite end-member (Fig. 6e), mainly at their rims transformed into Bt₃+Pl₃. A few specks of Ms₃ (Figs. 4f and 6e) isolated among Bt₃ and Pl₃ seem to be re-equilibrated relicts of Ph₂. Ilm₂ initially in contact with Ph_2 is also rimmed by concentric coronas of Pl_3 and Bt_3 (2 in Fig. 4f), and would have operated as a minor reactant, supplying some Ti (and Fe) to the newly formed biotite, whose high Ti-content is likely related to HT metamorphic conditions (*e.g.*, Henry and Guidotti 2002).

485 The stoichiometric balance of the phengite breakdown reaction was made by the 486 least-squares method (sample T138; for 1 mol of Ph_2):

487 1 Ph₂ + 3.79 Qz₂ + 0.20 Ilm₂ \rightarrow 0.21 Bt₃ + 1.25 Pl₃ + 2.01 Kfs₃ + 1.57 H₂O + 1 *R*,

488 where *R* is the vector of the residuals (*R*: $Si_{0.00}Al_{-0.48}Cr_{+0.00}Ti_{-0.23}Fe_{+0.23}Mn$. 489 $_{0.00}Mg_{+0.07}Ca_{+0.42}Na_{+0.80}K_{+0.47}O_{0.00}$). The conversion into volumes yields:

490 1 $Ph_2 + 0.321 Qz_2 + 0.022 Ilm_2 \rightarrow 0.229 Bt_3 + 0.446 Pl_3 + 0.774 Kfs_3$.

491 The minerals are correctly distributed between reactants and products, in volumetric 492 proportions comparable to what is observed (Fig. 4f). The increase in volume during the 493 reaction ($\Delta V_s/V_s = +0.079$) is also consistent with an evolution at decreasing pressure. 494 However, associated residuals are abnormally important, indicating that some additional 495 exchange took place during the reaction, which therefore occurred in an open system with 496 respect to the reaction domain considered.

497 Indeed, such an exchange could be linked to a mobile melt. Fluid-absent melting of 498 phengite typically occurs during the decompression of HP metapelitic and felsic rocks 499 (e.g., Vielzeuf and Holloway 1988; Hermann and Green 2001; Schmidt et al. 2004; 500 Hermann et al. 2006; Massonne 2009; Dokukina et al. 2017). Phengite melting in the 501 studied rocks is consistent with the calculated P-T pseudosections, where the phengite-502 out and solidus curves almost coincide (Fig. 7b-c). During the retrograde evolution, the 503 metapelites followed a *P*-*T* path sub-parallel to these two curves or, more likely, these 504 curves were shifted in the vicinity of the P-T path, by extraction of melt, as discussed 505 before (Section 6.2 iii). This evolution can explain the zonation observed at the rims of 506 both garnet porphyroblasts (decrease in X_{Fe} and X_{Grs} : Fig. 6a-d) and phengite flakes 507 (decrease in the celadonite end-member: Fig. 6e), which are consistent with the 508 decreasing values of x(Grt), z(Grt) and Si(Ph) predicted by the pseudosections during a 509 drop of *P* (Fig. 7b-c).

The final step of decompression and cooling is manifested by the formation of M_3 fine-grained aggregates, indicating an evolution into the melt-free stability fields of sillimanite and biotite (Fig. 7). Despite the tight spacing of isopleths in this region of the pseudosections, the compositional heterogeneity of the M_3 domains and minerals, which likely did not fully reach equilibrium, makes it difficult to specify the *P*-*T* conditions of this late metamorphic stage. Garnets in contact with the matrix, either "outer rims" of porphyroblasts or Grt₃, have the lowest grossular content [*z*(*Grt₃*): 0.13–0.05 for sample

517 T78; 0.10–0.08 for T52] and the highest X_{Fe} ratio [$x(Grt_3)$: 0.85–0.91 for T78; 0.87–0.93 518 for T52]. The X_{Fe} isopleth values for Bt₃ are: $x(Bt_3) = 0.58-0.62$ for T78; 0.67-0.69 for 519 T52. The isopleths do not intersect at a point but in a P-T region that provides a 520 reasonable estimate for the conditions of this stage (blue ellipses in Fig. 7b-d): 725-521 740°C at 7-8 kbar (T78); 720-730°C at 7.0-7.5 kbar (T52). Since equilibrium is 522 imperfectly achieved at the scale of the sample, we modeled a small homogeneous matrix 523 domain, similar to that of Figure 3g, for sample T52 (Fig. 7d). The compositions of the minerals [x(Grt₃): 0.87–0.88; z(Grt₃): 0.05–0.06; x(St₃): 0.89–0.94; x(Bt₃): 0.62–0.71] 524 suggest *P-T* conditions of 5.5–6.0 kbar, 675–725°C (blue ellipse in Fig. 7d) on a regular 525 geothermal gradient of ~15°C.km⁻¹. Moreover, modeling this microdomain reveals 526 527 features not shown by the bulk-rock pseudosection, like the local stability of St₃ and the 528 late increase of X_{Fe} in the garnet "outer rim" (see x(Grt) isopleths in Fig. 7d).

529 6.4. Petrological and geological implications

530 The metapelites studied here are distinctive regarding three particular circumstances 531 that, combined, make these rocks a kind of "geological miracle":

532 (i) *Eclogite-facies metapelites*. Eclogite-facies mafic metabasites, *i.e.*, true eclogites, 533 have been known for two centuries, but metapelites related to eclogites went unnoticed 534 for a long time, to the point that, in the 1980-90s, a controversy "foreign" vs. "in-situ" 535 raised the question of the relationships between eclogites and their metapelitic host, in 536 which traces of such metamorphism were not observed (see the historical summary in 537 Godard 2001). It has since been recognized that, among eclogite-facies rocks, metapelites 538 are not less rare than metabasites, but have rarely preserved their eclogite-facies imprint 539 due to the combined effects of retrograde metamorphism, rehydration and late 540 deformation. In this regard, the difficulty in specifying the P-T conditions prevailing at 541 the peak of metamorphism (Section 6.2) is usual for this type of rock. It remains that it is 542 amazing to observe in the Egéré terrane such rocks on a regional scale. Without wishing 543 to reignite the controversy, it should be noted that the mafic eclogites and eclogite-facies 544 metapelites from Egéré seem to have recorded slightly different P-T evolutions: In 545 Tighsi, 30–40 km N of the investigated Izzilatène area, the apparent *P*-*T* path of mafic 546 eclogites (19.6 kbar/695°C \rightarrow 15.5 kbar/775°C \rightarrow 9.3 kbar/695°C) does not exactly 547 superimpose on that of the host metapelites (17–19 kbar/850°C \rightarrow 9 kbar/750 °C) 548 (Doukkari et al. 2014; Arab et al. 2015). The same is observed in Izzilatène, between the 549 mafic eclogites (13–14 kbar/580°C \rightarrow 19 kbar/650–700 °C \rightarrow 8–9 kbar/700-750 °C: 550 Doukkari et al. 2015) and the investigated eclogite-facies metapelites (6.0-6.5 kbar/600-551 $630^{\circ}\text{C} \rightarrow 14-20 \text{ kbar}/800-820^{\circ}\text{C} \rightarrow 5.5-6.0 \text{ kbar}/675-725^{\circ}\text{C}$: this study). In both cases, 552 the apparent P peak is slightly lower, or at least less well defined, for the metapelites than 553 for the metabasites. On the other hand, the T peak seems higher by a hundred degrees for 554 metapelites compared with the mafic eclogite lenses that they contain. It is difficult to 555 imagine that these differences are linked to some major geological process. It makes more sense to invoke a bias linked to thermodynamic modeling, or even more surely a contrasting behavior of the two types of rock, since, relative to the metapelites, the metabasites preserve more easily the imprint of the P peak but record less easily the Tpeak because of their refractory properties.

- 560 (ii) *Nanogranitoids in eclogite-facies rocks*. Partial melting of eclogite-facies rocks is 561 still a matter of debate, not because melting under (ultra)high-pressure conditions is not 562 predicted by experiments and thermodynamics (e.g., Hermann et al. 2006), but because of the lack of irrefutable petrological evidence, especially for mafic eclogites. There is a 563 564 score of microstructures that suggest partial melting in some eclogite-facies rocks (see 565 review in Section 4), but a compelling evidence like melt inclusions sheltered in garnet and crystallized as "nanogranitoids" is rare. Such inclusions are well known in HT 566 567 granulite-facies migmatites (e.g., Cesare et al. 2009, 2015; Ferrero et al. 2012; Bartoli et 568 al. 2016), less frequently in HP granulite-facies rocks (Ferrero et al. 2018), and more 569 rarely in eclogite-facies quartzo-feldspathic rocks (Ferrero et al. 2015). This study 570 confirms that they exist under eclogite-facies conditions, despite less favorable 571 geotherms, although no evidence of melting has been found in the associated mafic 572 eclogites, likely because of their refractory composition.
- 573 (iii) Eclogite-facies metamorphism in a Neoproterozoic orogen. Recent studies (e.g., Doukkari et al. 2014, 2015; Arab et al. 2015; Arab 2016; and references therein) suggest 574 575 that the Egéré supracrustal sequence deposited in a continental rift/passive margin 576 environment, at an age of \geq 745 Ma. It has been suggested that these sediments were 577 intruded by mafic rocks and subducted deeply during the Neoproterozoic evolution of the 578 Hoggar, likely during the Upper Cryogenian-Lower Ediacarian (Liégeois et al. 2003; 579 Arab 2016), thus recording eclogite-facies conditions, before being exhumed. The LA-580 ICP-MS U-Pb analysis of a zircon megacryst from a neighboring eclogite-facies 581 metapelite, crystallized from the anatectic melt at conditions close to the T peak, gave an 582 Upper Cryogenian age of 654 ± 5 Ma (Bruguier et al. 2014). The Egéré terrane would 583 then represent an old passive margin that underwent a subduction-exhumation cycle 584 related to the Pan-African orogeny that led to the Western Gondwana assembly (Liégeois 585 et al. 2013; Berger et al. 2014; Liégeois 2019). The existence in the Hoggar of numerous 586 traces of an eclogite-facies metamorphism dating back to the Neoproterozoic is 587 remarkable, since the first true eclogites on Earth formed around 2.1 Ga ago (e.g., Loose 588 and Shenk 2018), indicating the emergence of modern plate tectonics. This type of 589 metamorphism remained quite rare in the Neoproterozoic, before gradually increasing 590 from the Paleozoic to the Present. The Egéré terrane belongs to the West Gondwana 591 Orogen (WGO), which stretches from Hoggar to the northeast of Brazil and is 592 characterized by a Cryogenian-Ediacarian subduction, which Ganade de Araujo et al. 593 (2014) consider as the earliest evidence of large-scale deep continental subduction and 594 Himalayan-scale mountains in the geological record.

595 **7. IMPLICATIONS**

596 The studied migmatitic metapelites preserve mineral zoning and textures related to 597 various stages of their prograde and retrograde metamorphic evolution, and provide 598 constraints on partial melting reactions under HP conditions. The following conclusions 599 can be made:

1. The metasedimentary Egéré unit (LATEA terrane, Central Hoggar, Algeria)
 mainly consists of kyanite-bearing metapelites, quartzites, marbles and mafic rocks that
 have undergone eclogite-facies metamorphism.

603 2. The studied metapelites consist of quartz, garnet, phengite, kyanite, rutile, with 604 secondary biotite, staurolite, muscovite, K-feldspar, plagioclase, sillimanite and ilmenite. 605 Garnet porphyroblasts show smoothed bell-shaped zoning patterns, with spessartine and 606 grossular decreases from core to rim. They grew during prograde metamorphism, 607 enclosing quartz, biotite, phengite, staurolite, kyanite, rutile and ilmenite. Coarse-grained 608 minerals, namely phengite, quartz and kyanite, form a foliated matrix and, together with 609 the rims of euhedral garnet porphyroblasts, constitute the peak paragenesis. Fine-grained 610 matrix minerals, mostly unoriented, like biotite, plagioclase, K-feldspar, muscovite, 611 staurolite and sillimanite, developed during exhumation throughout retrograde reactions 612 that produced a variety of intergrowths, coronas and pseudomorphs.

613 3. Two types of melting phenomena have occurred in the studied rocks: (i) H_2O -614 saturated melting during the prograde stage; and (ii) H_2O -unsaturated melting near the 615 peak and during decompression, likely associated with the dehydration melting of 616 phengite.

617 4. Polycrystalline micro-inclusions in garnet, with negative crystal shapes, formed 618 after entrapment of former melt and are therefore "nanogranitoids". Together with quartz 619 inclusions surrounded by feldspar, they provide evidence for anatexis during the prograde 620 evolution. Anatexis close to the metamorphic T peak gave a migmatitic structure to the 621 rocks. Biotite-plagioclase-bearing pseudomorphs are linked to dehydration melting of 622 phengite during exhumation.

623 5. A clockwise P-T path is suggested for the Egéré metapelites, based on micro-624 textures and P-T pseudosections. The recorded starting conditions of the prograde 625 evolution, evidenced from the mineral inclusions preserved in garnet, are 600–630°C and 626 6.0–6.5 kbar. The inferred peak assemblage (quartz, garnet, kyanite, phengite, rutile, 627 melt, \pm plagioclase, \pm K-feldspar) indicates a P peak of metamorphism at 14–20 kbar and a 628 T peak at 800–820°C. The assemblage garnet–biotite–staurolite–sillimanite–plagioclase– 629 K-feldspar–ilmenite, which occurs as late intergrowths, coronas and pseudomorphs, 630 formed at the expense of the HP paragenesis and documents the completion of the 631 retrograde evolution at ~6 kbar and 600-620°C.

6. The clockwise P-T path recorded by the Egéré HP metapelites is related to 633 continental subduction followed by exhumation during the Pan-African orogeny that led 634 to the amalgamation of the Hoggar shield. Recent geochronological data indicate that the 635 HP event likely took place at ~650 Ma.

7. The rocks studied are remarkable because (i) they are relatively well-preserved
eclogite-facies metapelites, (ii) they contain "nanogranitoids", rarely described so far in
eclogite-facies metamorphic rocks, and (iii) they indicate a subduction-related
metamorphism during a Neoproterozoic orogeny.

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928 **FIGURE CAPTIONS**

Figure 1. Geological context. (a) Geological sketch map of the Tuareg Shield (Black
et al. 1994; Liégeois 2019). (b) Geological map of the LATEA metacraton (after Liégeois
et al. 2003), showing the area studied. Ao: Aouilène; As-Is: Assodé-Issalane; Az: Azrou
n'Fad; Eg-Al: Egéré-Aleksod; La: Laouni; Se: Serouenout; Te: Tefedest; Tz: Tazat;
HKCA: high-potassium calc-alkaline rocks.

Figure 2. Outcrop photographs. (a) Upper metasedimentary unit of Egéré showing metapelites interbedded with marbles, quartzites and eclogite lenses (hills on the horizon). (b-d) Close-up views of the migmatitic metapelites studied, showing the alternation of various lithologies, in particular of leucosomes and melanosomes rich in garnet porphyroblasts, kyanite and phengite (d).

939 Figure 3. Mineral assemblages and microtextures of kyanite-bearing metapelites. 940 Plane-polarized (b-e, g, h) and cross-polarized (a, f) lights. (a) Garnet megacryst, rich in 941 inclusions defining a straight internal foliation and corroded by aggregates of biotite + 942 plagioclase (Bt₃+Pl₃). (b-d) Close-up views of various inclusions (Bt₁, Ilm₁, Qz₁, St₁, Rt₁) 943 in garnet porphyroblasts (Grt₁), some of which are surrounded by a plagioclase moat (Pl 944 in c). (e) Garnet + quartz skeletal intergrowth. (f) Phengite flake (Ph₂), partially resorbed 945 by aggregates consisting of biotite (Bt_3), plagioclase (Pl_3) and K-feldspar (Kfs₃). (g) 946 Symplectite mainly consisting of Bt₃, Pl₃, Kfs₃, Sil₃ and St₃, developed at the expense of 947 the peak paragenesis, in particular Grt_2 , Ph_2 , Ky_2 and Qz_2 . (h) Second generation of fine-948 grained staurolite (St₃) associated with late Bt₃, Kfs₃ and Sil₃.

949 Figure 4. Reaction microstructures in kyanite metapelites. BSE images (a-e), and 950 RGB image (f), the red, green and blue channels of which are the first three main 951 components of the PCA of element maps. (a) The HP paragenesis (M_2) , consisting of 952 garnet (Grt_{1-2}), phengite (Ph_2), kyanite (Ky_2) and quartz, is partially replaced by late M_3 953 assemblages (Bt_3+Pl_3). (b) Garnet megacryst (Grt_{1-2}) corroded by Bt_3+Pl_3 aggregates. (c) 954 Close-up view of quartz inclusions (Qz_1) in a garnet porphyroblast (Grt_1) , which are 955 surrounded by thin films of plagioclase (Pl). (d) Aggregates of Bt₃, Pl₃, Sil₃, St₃ and Ms₃, 956 having replaced the M₂ paragenesis ($Ph_2+Ky_2+Grt_2+Qz_2$); K-feldspar Kfs₃ mainly 957 developed as coronas around large Qz_2 grains. (e, f) Phengite flake (Ph₂) partially 958 resorbed by aggregates consisting of Bt₃, Pl₃, Ms₃ and Kfs₃, which forms a continuous 959 corona at contacts with Qz_2 ; lamellae of Ilm_3 and Bt_3 are exsolved parallel to the (001) 960 plane of phengite Ph₂; 1: numerous micro-inclusions of ilmenite inside muscovite (*i.e.*, 961 Ms₃+Ilm₃) after Ph₂, which was probably richer in Ti near Ilm₂; 2: Pl₃+Ilm₃ symplectite 962 between Ilm₂ and Ph₂.

Figure 5. Nanogranitoids in garnet porphyroblasts. Plane-polarized light (a) and BSE (b-f) images (sample T100). (a) Garnet megacrysts in a matrix with leucosomes and micaceous segregations. (b) Very small melt inclusions hosted in a garnet porphyroblast (Grt₁) and now crystallized as "nanogranitoids". (c-f) Close-up views of "nanogranitoids", which generally display the negative euhedral forms of the host garnet (d-f) and consist of quartz + biotite + plagioclase \pm K-feldspar \pm zircon.

969 Figure 6. Mineral compositions of the metapelites. (a-d) Composition of garnet in 970 samples T78 (a-b) and T138 (c-d). The profiles across garnet megacrysts (a, c) show a 971 smoothed compositional zoning at their cores, where a bell-shaped Mn profile is 972 observed; the zoning at the rims of profile (a) is similar to that of the inner rims of profile (b), whose outer rims show a reverse zoning when at contact with Bt₃+Pl₃ symplectites. 973 (e) White mica compositions in the diagram (Si+Fe²⁺+Mg) vs. (Al^{IV}+Al^{VI}), showing the 974 phengite substitution $(Si_{+1}[Fe^{2+},Mg]_{+1}+Al^{IV}_{-1}Al^{VI}_{-1})$. (f) Staurolite compositions in the 975 976 diagram X_{Mg} -Si. (g) Biotite compositions (diagram of Rieder et al. 1998). (h) Feldspar 977 compositions. The symbols and colors in (g) and (h) represent microstructural sites and978 samples, respectively (see the common legend).

Figure 7. *P*-*T* pseudosections for samples T78 and T52. Notations used for the isopleths: x(Grt/Bt/St) = (Fe/(Fe + Mg)); z(Grt) = (Ca/(Ca + Fe + Mg + Mn)); Si(Ph) = Si.The presumed prograde and retrograde *P*-*T* paths (red dashed curves) are constrained by various isopleths and isomodes; the blue dashed arrows indicate the shift of the solidus curve towards higher temperatures during melt extraction (see text).

















