

## Original Article

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# The onset of Neo-Tethys subduction in the Early Jurassic: evidence from the eclogites of the North Shahrekord Metamorphic Complex (Sanandaj-Sirjan Zone, W Iran)

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**Abstract**

Geodynamic models implying subduction of continental crust either consider this process happening during collision, when the continental margin of the lower plate attempts subduction, or in pre-collisional stages, when tectonic erosion of the upper plate or subduction of continental extensional allochthons drag continental crust in the subduction channel. In the Zagros orogen (W Iran), high-pressure rocks are known only from the Sanandaj-Sirjan Zone, NE of the Main Zagros Thrust. Here, eclogites of the North Shahrekord Metamorphic Complex suggest subduction of continental crust slices derived from the upper plate (Central Iran) during the onset of the Neo-Tethys subduction along the southern margin of Iran. Eclogites record a clockwise pressure-temperature-time path, with pre-eclogitic epidote-amphibolites-facies phase assemblages preserved in garnet cores, a high-pressure stage, and a subsequent retrogression at amphibolite-facies conditions. By means of forward thermodynamic modelling and  $^{40}\text{Ar}/^{39}\text{Ar}$  geochronology, the peak metamorphism has been constrained at 1.9–2.1 GPa and 550–600 °C, in the 191–194 Ma time span. The following retrogression during exhumation lasted at least until 144 Ma. Our data suggest that the onset of the Neo-Tethys subduction traces back prior to 190 Ma, involving together with the Neo-Tethys oceanic lithosphere also slices of the upper plate continental crust scraped off by means of tectonic erosion processes.

**1. Introduction**

High-Pressure (HP) metamorphic rocks are key to understand subduction-related processes (e.g. Agard *et al.* 2009; Cheng *et al.* 2015; Di Vincenzo *et al.* 2016). Eclogites can form at different times during the subduction-collision history (e.g. Spalla *et al.* 1996), from pre-collisional stages to final continent-continent collision, when continental crust of the lower plate can be subducted even at ultrahigh-pressure (UHP) conditions (e.g. Jolivet *et al.* 2003), thus providing invaluable information on the processes acting at mantle depths during the evolution of orogenic belts. Useful information on the tectonometamorphic evolution of eclogite-bearing units comes through the application of meso- and microstructural analyses, conventional geothermobarometry and forward thermodynamic modelling. Geochronological data add time constraints, framing the P-T evolution of high-pressure tectonometamorphic units in the whole geodynamic process that controls the birth and the evolution of an orogen.

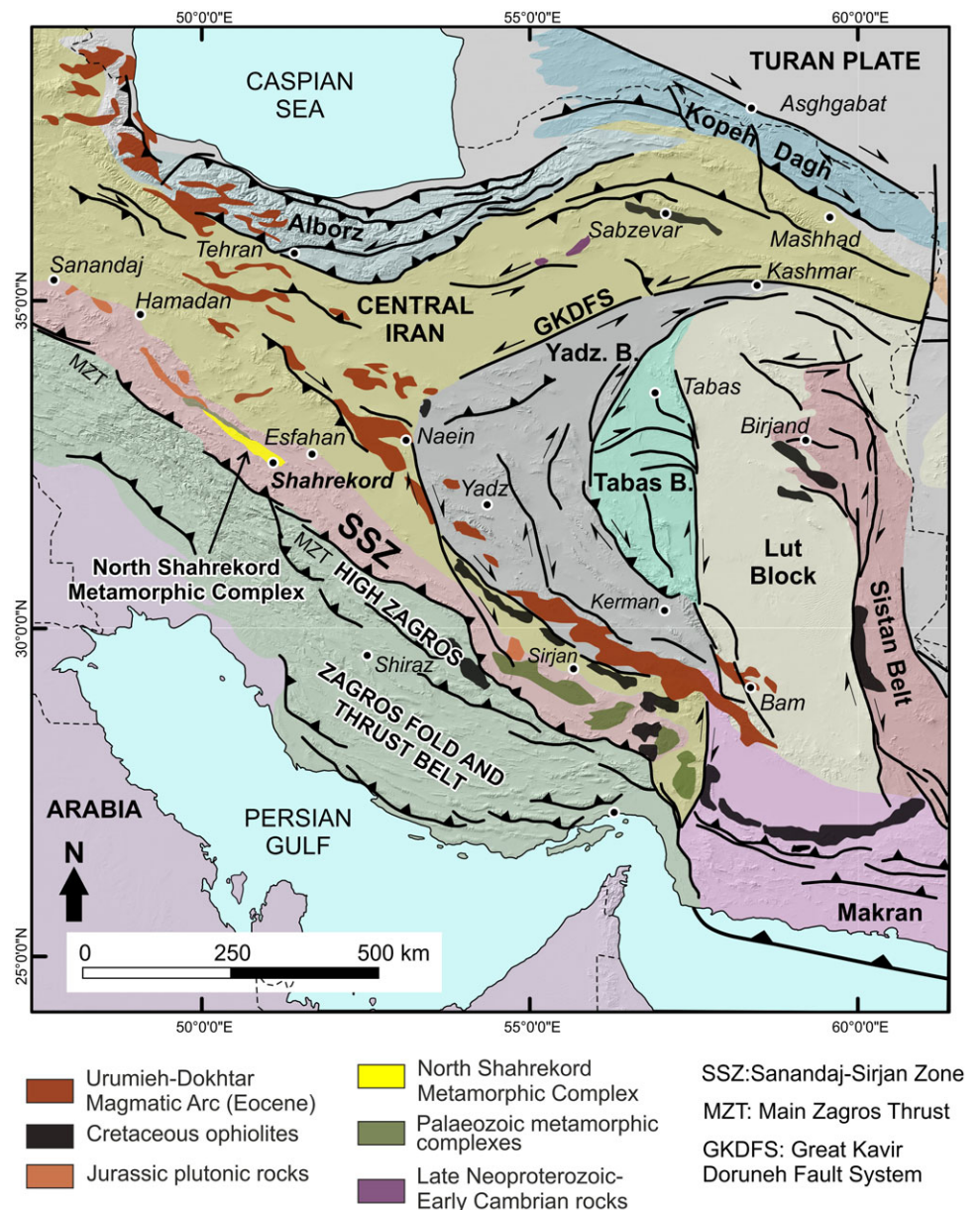
The  $^{40}\text{Ar}/^{39}\text{Ar}$  method is one of the most used dating techniques when facing metamorphic rocks because it well constrains their thermal history and exhumation information (Di Vincenzo & Palmeri, 2001; Warren *et al.* 2012). Given their stability at HP and ultra (U)HP conditions, amphiboles and phengitic white micas are usually ideal minerals for  $^{40}\text{Ar}/^{39}\text{Ar}$  dating.

The reconstructed P-T-t paths of continental HP rocks help to decipher the geodynamic scenario in which they formed, with special regard to their pre- or syn-collisional origin. Eclogites derived from continental crust may form at the end of the subduction of the intervening oceanic lithosphere between two continents, when slices of the continental crust of the lower plate are subducted below crustal depths. When derived from the upper plate, they can also form at the beginning or during the subduction in response to tectonic erosion processes (e.g. Polino *et al.* 1990; Massonne, 2012) that scrape off slices of the upper plate continental crust and drag them down along the subduction channel (e.g. Clift & Vannucchi, 2004; Vannucchi *et al.* 2008).

Here we present a multi-disciplinary study on the eclogites of the North Shahrekord Metamorphic Complex (NSMC, Sanandaj-Sirjan Zone, western Iran, Figure 1). The aim of this study is to unravel their Pressure-Temperature-time (P-T-t) path, providing information on the peak P-T conditions and their metamorphic evolution. The timing of the HP metamorphism

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and subsequent exhumation has been constrained by means of  $^{40}\text{Ar}/^{39}\text{Ar}$  geochronology on eclogites and amphibolites, their retrogressed counterparts. Reconstructing the evolution of the NSMC eclogites is a key step in the understanding of the geodynamic processes associated with the onset and later stages of the Neo-Tethys subduction in western Iran.

Our results provide new P-T-t constraints on the onset and duration of eclogite facies metamorphism and subsequent amphibolite-facies retrogression in the Sanandaj-Sirjan Zone (Figure 1). Furthermore, the new data presented here, together with previously published petrological, geochemical and geochronological data on eclogites- and amphibolite-facies rocks (e.g. Davoudian *et al.* 2008, 2016; Malek-Mahmoudi *et al.* 2017; Jamali Ashtiani *et al.* 2020; Maghdour-Mashhour *et al.* 2021; Wan *et al.* 2023), refine the geodynamic model for the evolution of the Sanandaj-Sirjan Zone in western Iran. We propose here a new scenario involving the subduction of continental crust induced by the tectonic erosion of the overriding plate.

## 2. Geological background

The Alpine-Himalayan orogenic belt resulted from the closure of the Neo-Tethys Ocean and consequent continental collision between the Eurasian and the African-Arabian-Indian plates. In the central part of the Neo-Tethys, the collision between Eurasia and Arabia and several continental microplates (i.e. the Cimmerian blocks) caused the formation of the Turkish-Iranian plateau and of the associated orogenic belts during the Mesozoic-Cenozoic (Brunet & Cloetingh, 2003). The long-lasting convergence between the Eurasian and Arabian plates promoted the formation of the Zagros orogen (Agard *et al.* 2011), which consists of several NW-SE trending tectonic units. These tectonic units from NE to SW are as follows: (i) the Urumieh-Dokhtar Magmatic Arc, an arc mainly composed of Cenozoic plutonic and volcanic rocks (Schröder, 1944; Förster, 1974; Berberian & King, 1981; Berberian *et al.* 1982; Alavi, 1994); (ii) the Sanandaj-Sirjan Zone (SSZ, see Figure 1 in Stöcklin, 1968), representing a partially independent magmatic and metamorphic belt exposed to the NE of

the Zagros orogen (Agard *et al.* 2005; Mohajjel & Fergusson, 2014; Azizi *et al.* 2015a, 2015b; Moghadam *et al.* 2015; Shakerardakani *et al.* 2015; Hassanzadeh & Wernicke, 2016); (iii) the High Zagros consisting of imbricated tectonic slices of Mesozoic limestones, radiolarites and remnants of obducted ophiolites (Agard *et al.* 2005); (iv) the Zagros Fold-and-Thrust Belt; and (v) the Mesopotamian-Persian Gulf foreland basin (Berberian & King, 1981; Alavi, 1994; Mohajjel & Fergusson, 2000). The Main Zagros Thrust (MZT) is identified as the suture zone between the Arabian plate and the Cimmerian blocks, with the SSZ that developed in the upper plate during the subduction of the Neo-Tethys Ocean.

The SSZ is located along the southwestern edge of the Iranian plateau (Figure 1), running parallel to the Zagros orogen for about 1500 km from eastern Anatolia to the north-western Makran area (McCall, 2002). The SSZ consists of late Neoproterozoic and Pan-African basement (Crawford, 1977; Hassanzadeh *et al.* 2008; Nutman *et al.* 2014) that discontinuously crops out beneath Palaeozoic to Neogene metamorphic, igneous and sedimentary units (Agard *et al.* 2011; Jamshidi Badr *et al.* 2013; Mohajjel *et al.* 2003; Mohajjel & Fergusson, 2014; Moghadam *et al.* 2015; Shakerardakani *et al.* 2015; Sheikholeslami, 2015). The Palaeozoic sedimentary and magmatic record (Alirezaei & Hassanzadeh, 2012; Moghadam *et al.* 2015) of the SSZ is related to the opening of the Neo-Tethys that was in turn related to the rifting of continental fragments (i.e. the Cimmerian block, Ricou, 1994; Stampfli & Borel, 2002; Şengör & Yilmaz, 1981; Şengör, 1984; Zanchi *et al.* 2015) away from the northern margin of Gondwana during the Permian (Alavi, 1994; Angiolini *et al.* 2013, 2015; Agard *et al.* 2011; Hassanzadeh & Wernicke, 2016; Mattei *et al.* 2015; Moghadam *et al.* 2015; Zanchi *et al.* 2009a). During the Cimmerian orogeny (Late Triassic), these crustal blocks collided with the Eurasian plate following the closure of the Palaeo-Tethys (e.g. Wilmsen *et al.* 2009; Zanchi *et al.* 2009a; Zanchetta *et al.* 2013a; Zanchi *et al.* 2016). Even if the evolution of the Cimmerian blocks after their collision is still not completely resolved (Zanchi *et al.* 2009b; Zanchi *et al.* 2021 and references therein), their northern margin runs along the Palaeo-Tethys suture in north Iran and extends to northern Afghanistan, Pakistan, Pamir, Tibet and China (Angiolini *et al.* 2013, 2015; Zanchetta *et al.* 2018), and their southern boundary lies along the Neo-Tethys suture, namely the Main Zagros Thrust (MZT).

The switch from a passive continental margin to a continental arc in the SSZ occurred after the Cimmerian orogenic event, during the Late Triassic-Early Jurassic (Hassanzadeh & Wernicke, 2016; Maghdour-Mashhour *et al.* 2021), when the Neo-Tethyan NNE-NE-dipping subduction likely started. The Mesozoic to Cenozoic convergence finally led to the collision between the Arabian and the Eurasian plates (Mohajjel *et al.* 2003; Agard *et al.* 2005; Ghasemi & Talbot, 2006; Moritz *et al.* 2006). From the onset of the Neo-Tethys subduction to the collision with Arabia, the SSZ was not only affected by subduction-related magmatism but also by deformative and metamorphic events, mainly of Jurassic age (e.g. Fergusson *et al.* 2016), recorded in the several metamorphic-magmatic complexes scattered all along the SSZ (Jamshidi Badr *et al.* 2013), the most of them showing greenschist to amphibolite and even eclogite facies conditions (Mohajjel & Fergusson, 2000; Sheikholeslami *et al.* 2008; Ghasemi & Poor Kermani, 2009; Davoudian *et al.* 2016). Following Sheikholeslami (2015), the Jurassic rocks of the SSZ have been affected by, at least, two syn-metamorphic regional deformation phases: the first occurring in the Early Jurassic includes four deformation events; the second one resulted in three deformation events affecting Lower Jurassic

turbidites. During the first deformation phase, the metamorphic peak was reached at c. 187–180 Ma (U-Pb dating on Zrn; Fazlnia, 2007), whereas the HT peak metamorphism during the second deformational phase was constrained at c. 162 Ma (Sadegh *et al.* 2020).

The study area is located in the central SSZ in the North Shahrekord Metamorphic Complex (NSMC) west of Esfahan, belonging to the complexly deformed sub-zone (Figures 1 and 2). Here rare eclogite bodies occur as lenses or small-sized boudins within ortho- and paragneisses derived from the Gondwanan Pan-African basement of Iran (Davoudian *et al.* 2008, 2016; Nutman *et al.* 2014). The Gondwanan origin of the SSZ metamorphic belt has been proven (Fergusson *et al.* 2016) based on zircon ages pertaining to the Pan-African basement, which is exposed in wide areas of Central and Northwestern Iran (e.g. Hassanzadeh *et al.* 2008; Rahmati-Ilkhchi *et al.* 2011). U-Pb zircon ages also confirmed the Pan-African palaeogeographic affinity of granitic and metamorphic rocks near Golpaygan and around Lake Urumieh in the north-western sector of the SSZ (Fergusson *et al.* 2016 with references). Pan-African basement ages have been also documented in the central and northern Iranian Cimmerian continental blocks (Fergusson *et al.* 2016).

Eclogites of the NSMC occur as metre- to decametre-sized boudins hosted in amphibolites and paragneisses. Other units forming the NSMC consist of low- to very low-grade rocks such as schist, phyllite, marble, slate and metamorphosed sandstone, limestone, shale and volcanic rocks, with a possible Triassic to Jurassic age of the protoliths (U-Pb dating on Zrn; Davoudian *et al.* 2006; Babaahmadi *et al.* 2012).

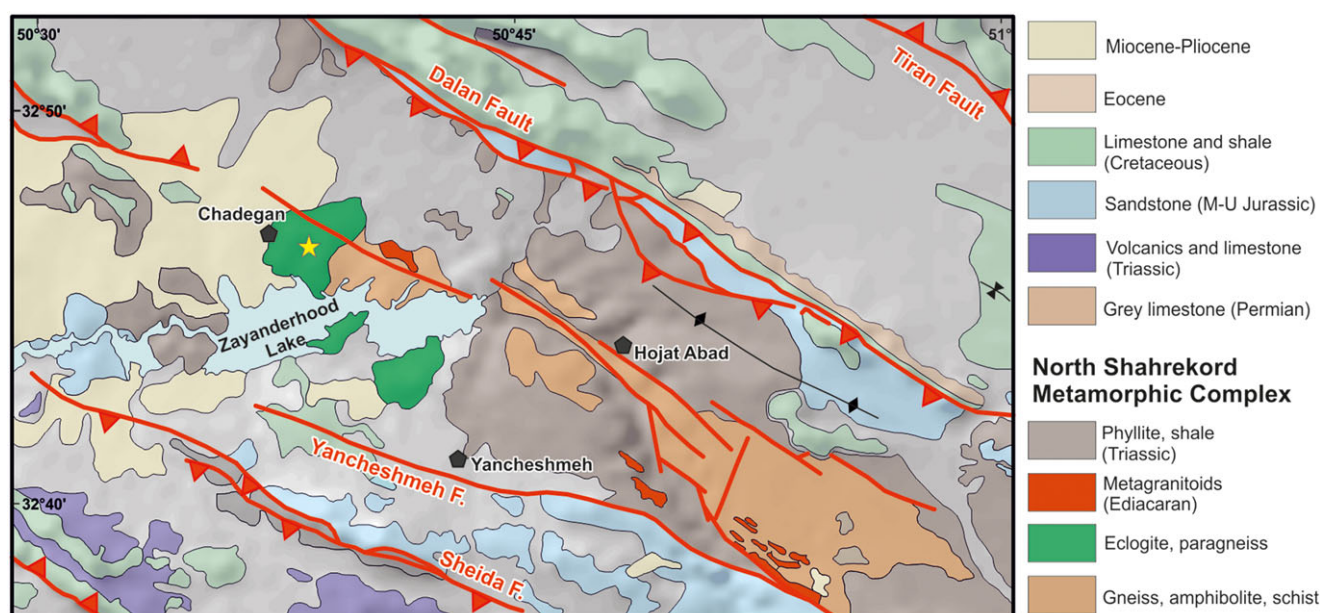
### 3. Methods

#### 3.a. Electron microprobe analyses

Electron microprobe analyses (EMPA) were carried out using a JEOL 8200 Super Probe EMP at the Department of Earth Sciences 'A. Desio', University of Milano. WDS (wavelength dispersive spectroscopy) quantitative chemical analyses and WDS X-ray elemental maps were performed on carbon-coated petrographic thin sections. Data acquisition was performed using an accelerating voltage of 15 kV, a beam current of 5 nA with a spot size of 3 µm on micas and 1 µm on other minerals. Oxides and natural silicates were used as standards. Analyses were recalculated to atom per formula unit (a.p.f.u.) as follows: garnet analyses were recalculated based on 8 cations and 24 charges; white mica was recalculated based on 11 oxygens and considering all Fe as Fe<sup>2+</sup>; clinopyroxene data were elaborated following the procedure by Cawthorn & Collerson (1974) based on 12 oxygens and considering Fe<sup>3+</sup> as the acmite component; amphiboles analyses were recalculated based on 23 oxygens and 13 cations plus K, Na, Ca. Amphibole data were classified and plotted according to Hawthorne & Oberti (2007).

#### 3.b. <sup>40</sup>Ar/<sup>39</sup>Ar geochronology

Samples selected for <sup>40</sup>Ar/<sup>39</sup>Ar dating were crushed and sieved in order to separate white mica (in the fraction 250–500 µm) and amphibole (in the fraction 125–250 µm) for stepwise heating experiments. The separates were enriched in white mica and amphibole using a Frantz magnetic separator and handpicked under a stereomicroscope. Mineral separates were then cleaned ultrasonically in deionised water and wrapped in aluminium foil. Samples and standards were irradiated for 30 MWh in a fast neutron flux at the McMaster University Research Reactor (Hamilton, CA).



**Figure 2.** (Colour online) Simplified geological map of the NSMC in the Zayanderhood Lake area according to our field surveys and after Davoudian *et al.* (2016). The intrusion age of the metagranitoid has been constrained to be Ediacaran (Jamali Ashtiani *et al.* 2020). The yellow star shows the location of Chadegan eclogites analysed in this study.

Stepwise heating experiments were performed at the ‘Laboratorio di Geocronologia’ of the University of Milano–Bicocca following the procedure reported in Montemagni & Villa (2021) and Montemagni & Zanchetta (2022). Sample and standards were loaded in a double vacuum resistance furnace attached to a NuInstruments™ Noblesse® noble gas mass spectrometer, equipped with one Faraday collector with a 1011  $\Omega$  resistor and two MasCom™ ion counters. The samples were heated in 10 steps for 20 min each at temperatures from 620 °C to 1300 °C. The irradiation intensity factor,  $J$ , was interpolated for each sample from the equation defined by the  $J$  values of the monitors. The  $^{40}\text{K}$  decay constant used for the age calculation was  $5.543 \times 10^{-4} \text{ Ma}^{-1}$  (Steiger & Jäger, 1977). To control the vertical flux gradient, monitors of McClure Mountain hornblende (MMhb) were interlayered with the sample disk wraps. The MMhb age was assumed to be  $523.98 \pm 0.12 \text{ Ma}$  (Schoene & Bowring, 2006). Data reduction procedures, corrected for mass spectrometer background, ion counter gains, blank measurements, source fractionation and decay of  $^{37}\text{Ar}$  since irradiation, were processed in the in-house CalcolAr Excel spreadsheet, in which also all the associated uncertainties are propagated.

### 3.c. Thermodynamic modelling

Thermodynamic modelling in the  $\text{SiO}_2$  (46.71 wt%),  $\text{Al}_2\text{O}_3$  (15.37 wt%),  $\text{FeO}$  (10.25 wt%),  $\text{MgO}$  (8.30 wt%),  $\text{CaO}$  (10.47 wt%),  $\text{Na}_2\text{O}$  (2.97 wt%),  $\text{K}_2\text{O}$  (0.87),  $\text{TiO}_2$  (1.41 wt%),  $\text{MnO}$  (0.18 wt%),  $\text{H}_2\text{O} + \text{CO}_2$  (1.80 wt%, LOI in Table 1) compositional system (sample Z5S in Table 1) was performed with the software package Perple\_X 7.1.5 (<http://www.perplex.ethz.ch>; Connolly, 2005), using the thermodynamic database of Holland & Powell (2011) (hp62ver.dat). The whole rock major elements composition was determined on the best-preserved eclogite (sample Z5S) by ICP-MS analysis at the ACME analytical laboratories (Vancouver, Canada) by Inductively Coupled Plasma Emission Spectrometry (ICP-ES) for major and minor elements and Inductively Coupled Plasma Mass Spectrometry (ICP-MS) for trace elements, including the rare earth elements, after fusion with lithium metaborate/tetraborate and

digestion by diluted  $\text{HNO}_3$ . Loss on ignition (LOI) was determined by weight difference after ignition at 1000 °C. Blank analyses were always below the minimum detection limit for each element, and the analytical protocol included the analysis of the reference materials (standards) OREAS184, SO-19, OREAS30A and OREAS262.

We used the following solution models: Gt(W), Chl(W) and Ctd(W) for garnet, chlorite and chloritoid from White *et al.* (2014), Omph(GHP) for clinopyroxene from Green *et al.* (2007), CAMph(G) for amphibole from Green *et al.* (2016), Pheng(HP) for white mica from Holland & Powell (1998), Sp(WPC) for spinel after White *et al.* (2002), feldspar from Fuhrman & Lindsley (1988) and cCcM(EF) for carbonate from Franzolin *et al.* (2011). The  $\text{H}_2\text{O}-\text{CO}_2$  (CORK) equation of state was taken from Holland & Powell (1998). Considering the low vol% amount of epidote (only in small inclusions in garnet) and of aegirine component in clinopyroxene, for the sake of simplicity, we considered the total iron in the bulk as  $\text{Fe}^{2+}$ .

### 4. Field occurrence and relationships

The NSMC (North Shahrekord Metamorphic Complex) consists of three main tectonometamorphic units (Davoudian *et al.* 2016): an eclogite-bearing unit located in the area of the Zayandeh-Rood Lake, a low-grade unit extending SE of the lake, mainly made of phyllites (Figure 2), and a third unit, mainly consisting of gneisses, amphibolites and schists, cropping out to the east in a tectonic window within the low-grade metamorphic unit. Lenses of weakly deformed metagranitoids of Ediacaran intrusion age (U-Pb on Zrn, Jamali Ashtiani *et al.* 2020) occur in the two units.

The contacts between the eclogite-bearing unit and surrounding ones are hardly visible due to poor exposures (Figure 3a). Amphibolites of the third unit frequently show a (proto)mylonitic fabric, suggesting that they are in contact with the rocks of the HP unit through a ductile shear zone. Eclogites occur as metre-sized to several decametre-sized boudins within country rocks (Figure 3c), which mainly consist of garnet-bearing paragneiss and granitoid

**Table 1.** Whole rock analyses of eclogites. Major elements are expressed as oxide wt%, trace elements as ppm

Sample	Z3S	Z4S	Z5S	Z7S
SiO <sub>2</sub>	45.77	48.41	46.71	46.82
TiO <sub>2</sub>	1.16	1.56	1.41	1.39
Al <sub>2</sub> O <sub>3</sub>	17.09	14.68	15.37	15.04
Fe <sub>2</sub> O <sub>3</sub>	10.37	12.56	11.39	11.89
MgO	9.01	7.29	8.30	8.11
CaO	11.25	10.87	10.47	10.55
Na <sub>2</sub> O	2.09	2.61	2.97	2.76
K <sub>2</sub> O	0.60	0.53	0.87	0.83
P <sub>2</sub> O <sub>5</sub>	0.09	0.13	0.14	0.09
MnO	0.16	0.20	0.18	0.18
Cr <sub>2</sub> O <sub>3</sub>	0.051	0.025	0.058	0.048
LOI	2.1	0.8	1.8	1.7
Sc	35	44	41	39
V	234	306	277	298
Cr	349	171	397	328
Co	48.6	44.2	43.4	42.1
Ni	135	45	90	48
Ga	16.0	17.3	16.6	15.6
Rb	15.2	20.6	20.6	19.8
Sr	211.9	246.8	184.4	201.5
Y	21.9	31.2	27.4	28.3
Zr	73.3	107.1	97.1	103.4
Nb	4.2	6.2	6.4	6.1
Ba	389	865	148	354
Cs	0.1	0.4	0.1	0.1
La	6.2	7.5	8.0	7.7
Ce	13.6	17.8	18.0	18.1
Pr	1.94	2.63	2.56	2.66
Nd	8.9	12.6	12.5	12.4
Sm	2.78	3.82	3.62	3.73
Eu	1.03	1.47	1.22	1.31
Gd	3.66	5.12	4.61	5.07
Tb	0.61	0.82	0.74	0.78
Dy	3.90	5.49	4.84	4.98
Ho	0.82	1.17	1.04	1.12
Er	2.34	3.29	3.04	3.12
Tm	0.37	0.48	0.42	0.41
Yb	2.23	2.98	2.74	2.78
Lu	0.34	0.46	0.41	0.42
Hf	2.1	3.1	2.4	2.6
Th	0.5	0.7	0.8	0.6
U	0.2	0.2	0.2	0.2

(Continued)

**Table 1.** (Continued)

Sample	Z3S	Z4S	Z5S	Z7S
Ta	0.3	0.4	0.4	0.3
ΣREE	48.72	65.63	63.74	64.58
ΣLREE	33.42	44.35	44.68	44.59
ΣHREE	14.27	19.81	17.84	18.68
Eu/Eu*	0.99	1.02	0.91	0.92
(La/Yb) <sub>n</sub>	1.994	1.805	2.094	1.987
(La/Sm) <sub>n</sub>	1.440	1.267	1.427	1.333
(Gd/Yb) <sub>n</sub>	1.358	1.421	1.392	1.509

Note: LOI: Loss On Ignition; n subscript – normalization to chondrite values (McDonough & Sun, 1995); ΣLREE – light rare earth element sum; ΣHREE – heavy rare earth elements sum; Eu/Eu\* = E<sub>un</sub>/RADQ(S<sub>mn</sub> × G<sub>dn</sub>).

orthogneiss (Figure 3d and 3e). The regional foliation in paragneiss and orthogneiss dips to S-SSW at low to medium angle (Figure 3). Detailed structural relationships within the eclogite-bearing unit of the NSMC are not available since the strong weathering and the dispersed occurrence of outcrops (Figure 3a) significantly hamper their reconstruction in the field. However, where observable, the eclogitic boudins are always hosted in garnet paragneiss (Figure 3c).

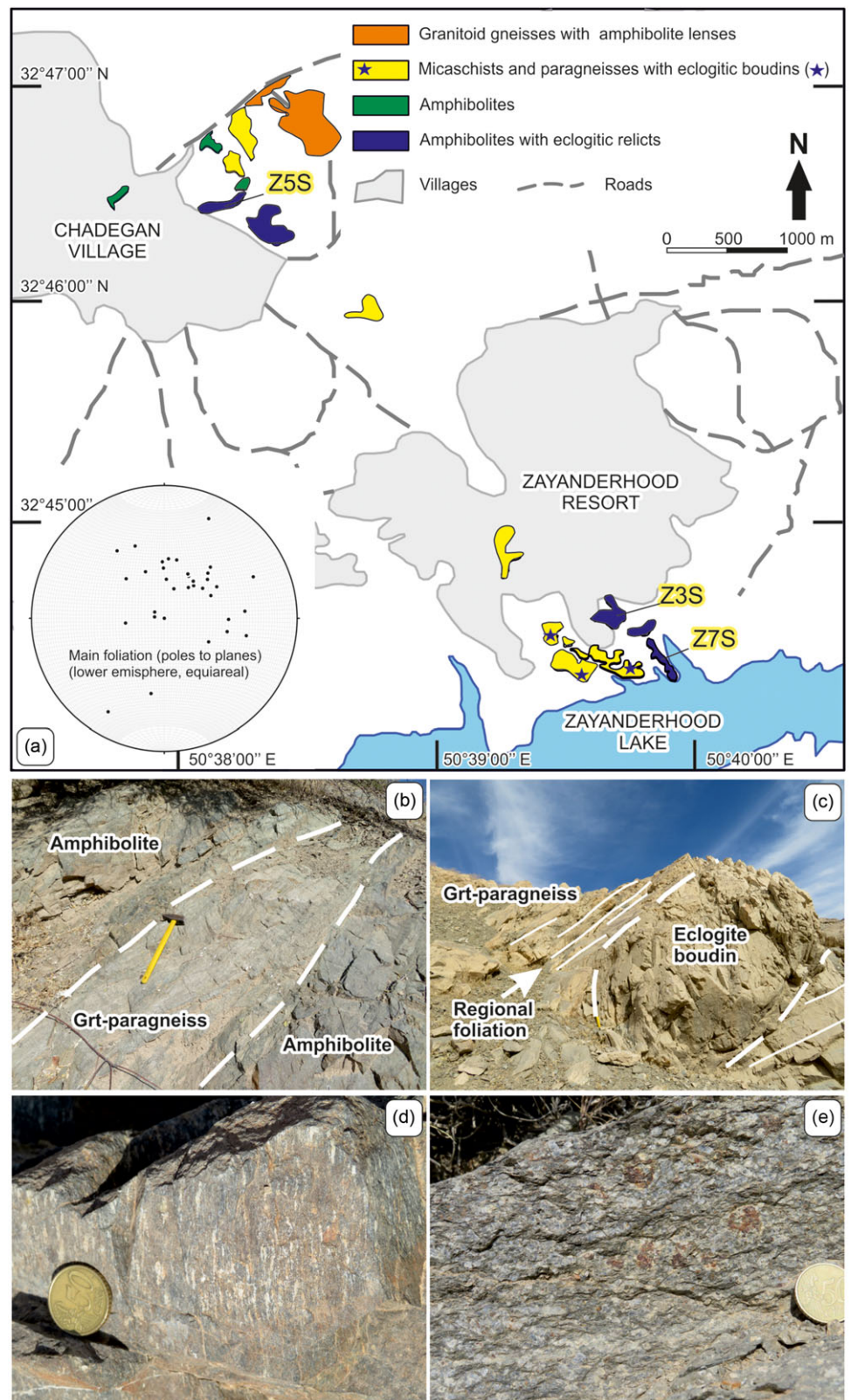
Eclogites display different degrees of retrogression, ranging from almost fresh eclogites with the HP mineral phase assemblage still well preserved, to garnet and epidote amphibolites. Retrogression of the eclogitic lenses is coupled with an increase in deformation, with former boudins that are progressively stretched along the main foliation of the hosting paragneisses (Figure 3b). Eclogitic lenses display a foliation and/or compositional layering developed at small angle with respect to the paragneiss foliation that wraps around the boudins. The mineralogical layering is defined by alternating omphacite-phengite rich and garnet-amphibole rich levels, with the shape preferred orientation of omphacite, phengite and clinozoisite that marks the foliation.

## 5. Metamorphic evolution of the NSMC eclogites

Petrographic observations were performed on all lithologies of the metamorphic complex. Representative microstructures of the main equilibrium phase assemblages are reported in Figure 4. Mineral abbreviations are after Whitney & Evans (2010) except for white mica (Wm).

Our study was focussed both on fresh or only partially retrogressed eclogite samples (Figure 4a-f), in order to reconstruct the evolution of the equilibrium phase assemblages and to define the P-T evolution of the eclogite-bearing unit of the NSMC. The degree of retrogression after the HP stage is variable from eclogites still almost completely preserving the HP phase assemblage (Figure 4a-b), to eclogites completely retrogressed into epidote amphibolites.

Two-mica orthogneiss (Figure 4g) and garnet-bearing paragneiss (Figure 4h) are also found in the NSMC, either showing a foliation made by micas enveloping feldspar (Figure 4g) or garnet (Figure 4h) porphyroblasts. Most of the orthogneisses were garnet-bearing, as testified by the occurrence of abundant chlorite pseudomorphs after garnet porphyroblasts (Figure 4g).

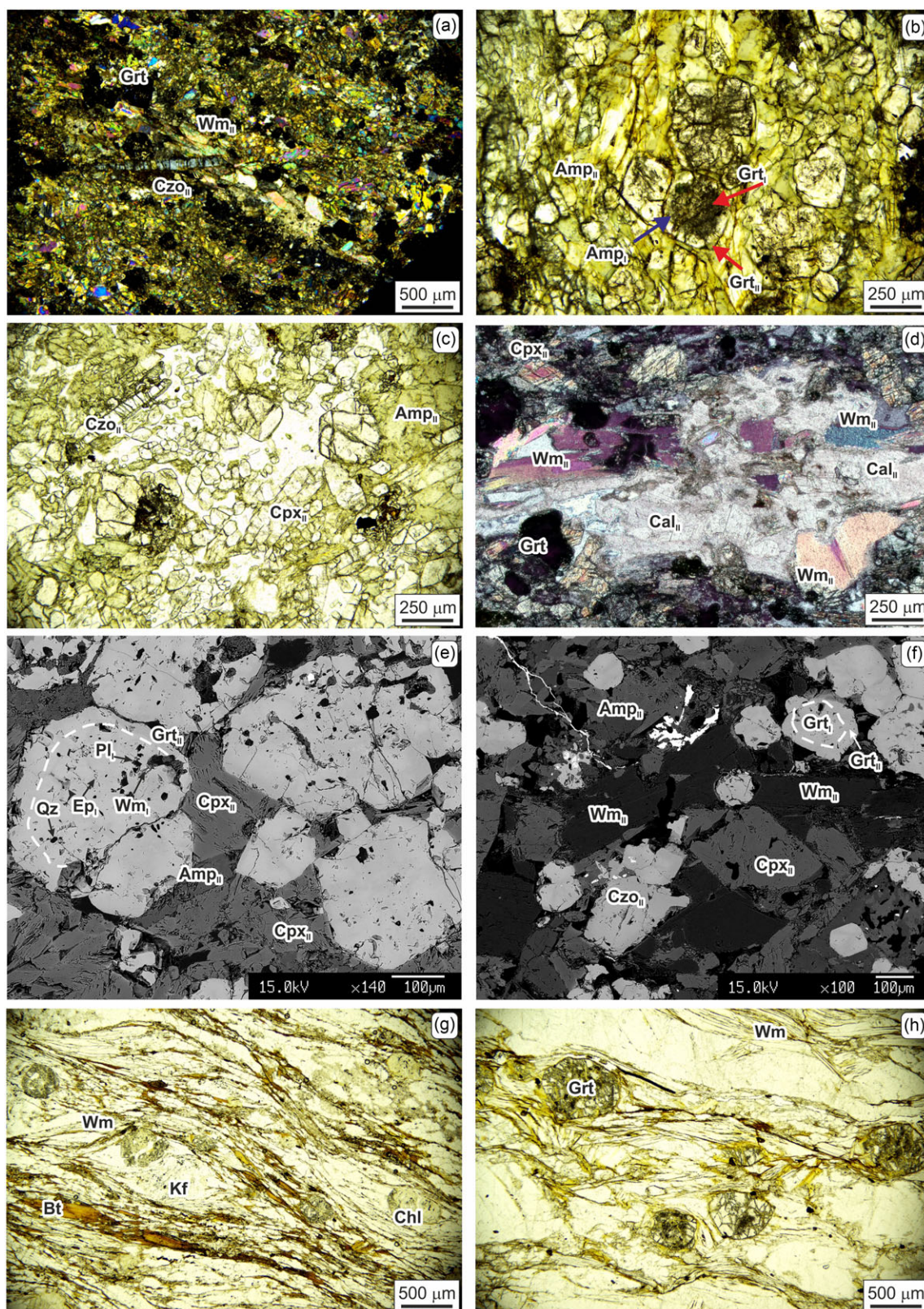


**Figure 3.** (Colour online) (a) Outcrops map of the NSMC rocks in the Chadegan area according to our field surveys. The location of the analysed samples has been reported. Field aspect of eclogites and surrounding rocks: (b) amphibolite layers stretched along the main foliation of the hosting paragneisses; (c) eclogitic boudins hosted in garnet paragneiss. The regional foliation in paragneiss has been highlighted. (d) Eclogite texture as appearing in the field, with a marked foliation and weakly developed compositional layering; (e) Grt-bearing granitoid orthogneisses.

The preserved eclogites (Figure 4a) are made of omphacite, garnet, white mica, clinozoisite, amphibole, epidote, plagioclase and quartz; rutile and ilmenite are the main accessory phases, whereas apatite and zircon are rare. Calcite also occurs in some samples, in lath-shaped crystals parallel to the omphacite foliation.

Based on petrographic and microstructural observations, three equilibrium phase assemblages, and relative metamorphic stages, can be reconstructed and summarized as follows (Figure 5):

(1) pre-eclogite facies preserved as inclusions within garnet cores (stage I in Figure 5); (2) a HP eclogite facies stage (stage II in



**Figure 4.** (Colour online) Microphotographs of rock fabrics. (a) The preserved eclogites showing an inequigranular texture, with garnet porphyroblasts of 5–12 mm embedded in a matrix; (b) Garnet porphyroblasts showing inclusions-rich core (Grt<sub>I</sub>) and clear rim (Grt<sub>II</sub>); (c) Clinozoisite (Czo<sub>II</sub>) occurs along the HP foliation in retrogressed eclogite; (d) Cal<sub>II</sub> laths aligned parallel to the eclogitic foliation marked by Wm<sub>II</sub>. Two-mica orthogneisses (g) and garnet-bearing paragneisses (h) showing a foliation made by micas enveloping feldspar (g) or garnet (h) porphyroblasts. (e) Garnet porphyroblasts preserving in their cores minerals of stage I (i.e. pre-eclogitic); (f) Stage II phase assemblage with Amp<sub>II</sub> + Wm<sub>II</sub> + Cpx<sub>II</sub> + Grt<sub>II</sub> + Czo<sub>II</sub>.

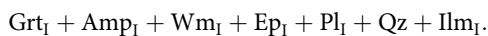
	Stage I Pre-eclogite	Stage II Eclogite	Stage III Amphibolite
Cpx		●●●●●●●●	●●●●●●●●
Grt	●●●●●●●●	●●●●●●●●	
Ep	●●●●●●●●		●●●●●●●●
(C)zo		●●●●●●●●	●●●●●●●●
Amp	●●●●●●●●	●●●●●●●●	●●●●●●●●
Wm	●●●●●●●●	●●●●●●●●	
Pl	●●●●●●●●		●●●●●●●●
Qz	●●●●●●●●	●●●●●●●●	●●●●●●●●
Cal		●●●●●●●●	
Rt		●●●●●●●●	
Ilm	●●●●●●●●		

**Figure 5.** Reconstructed textural equilibrium phase assemblages derived from microstructural observation for pre-eclogite (I), eclogite (II) and amphibolite (III) metamorphic evolution stages.

Figure 5) and (3) a late amphibolite-facies retrogression (stage III in Figure 5). Subscripts of mineral labels refer to the metamorphic stages in which they formed and/or were re-equilibrated.

### 5.a. Pre-eclogite facies stage (I)

Due to the strong overprint during subsequent deformation and metamorphic recrystallization, most of the mineral assemblages and textures belonging to the pre-eclogite facies metamorphism have been preserved only as inclusions in the cores of garnet porphyroblasts (Figure 5b). The preserved phase assemblage (Figure 5) is represented by:

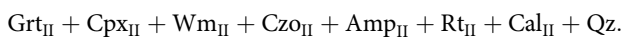


### 5.b. HP eclogite facies stage (II)

The eclogites display an inequigranular texture, with garnet porphyroblasts of 5–12 mm embedded in a matrix mainly composed of omphacite, amphibole and zoisite. Garnets have frequently an inclusions-rich core ( $\text{Grt}_I$ ) and a clear rim ( $\text{Grt}_{II}$ ).

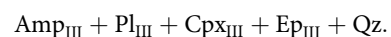
The rim of garnet porphyroblasts ( $\text{Grt}_{II}$ ) appears pre- to syn-kinematic with respect to the HP foliation (Figure 4b), which is individuated by the shape preferred orientation of omphacite ( $\text{Cpx}_{II}$ ), amphibole ( $\text{Amp}_{II}$ ) and white mica ( $\text{Wm}_{II}$ ). Clinzoisite ( $\text{Czo}_{II}$ ) also occurs along the HP foliation in some samples (Figure 4c) or crosscut the foliation ( $\text{Czo}_{III}$ ). Calcite also occurs, in lath-shaped crystals parallel to the omphacite and white mica foliation (Figure 4d), likely derived from previous aragonite at HP and subsequently transformed into calcite during decompression (Dey *et al.* 2023). Tiny rutile crystals ( $\text{Rt}_{II}$ ) occur mainly close to  $\text{Amp}_{II}$  or along omphacite-garnet interfaces.

The HP eclogite facies were constrained by the phase assemblage (Figure 5):



### 5.c. Amphibolite-facies stage (III)

The retrogression under amphibolites-facies conditions results in the partial substitution of omphacite ( $\text{Cpx}_{II}$ ) by pseudomorphic aggregates of amphibole ( $\text{Amp}_{III}$ ) + plagioclase ( $\text{Pl}_{III}$ ) or diopside ( $\text{Cpx}_{III}$ ) + plagioclase ( $\text{Pl}_{III}$ ), which is evidence of the late partial breakdown of omphacite during water-present decompression after HP metamorphism (O'Brien & Rötzler, 2003; Martin, 2019). Extremely fine-grained diopside + hornblende + plagioclase symplectites grew after the HP Ca-Na-clinopyroxene ( $\text{Cpx}_{II}$ ). The amphibole formed at this stage not only occurs in symplectites but also crystallizes in the matrix, partially substituting large  $\text{Amp}_{II}$  crystals with prismatic habit that were likely aligned along the HP foliation. Epidote ( $\text{Ep}_{III}$ ) has been also observed in some samples. The phase assemblage (Figure 5) is represented by:

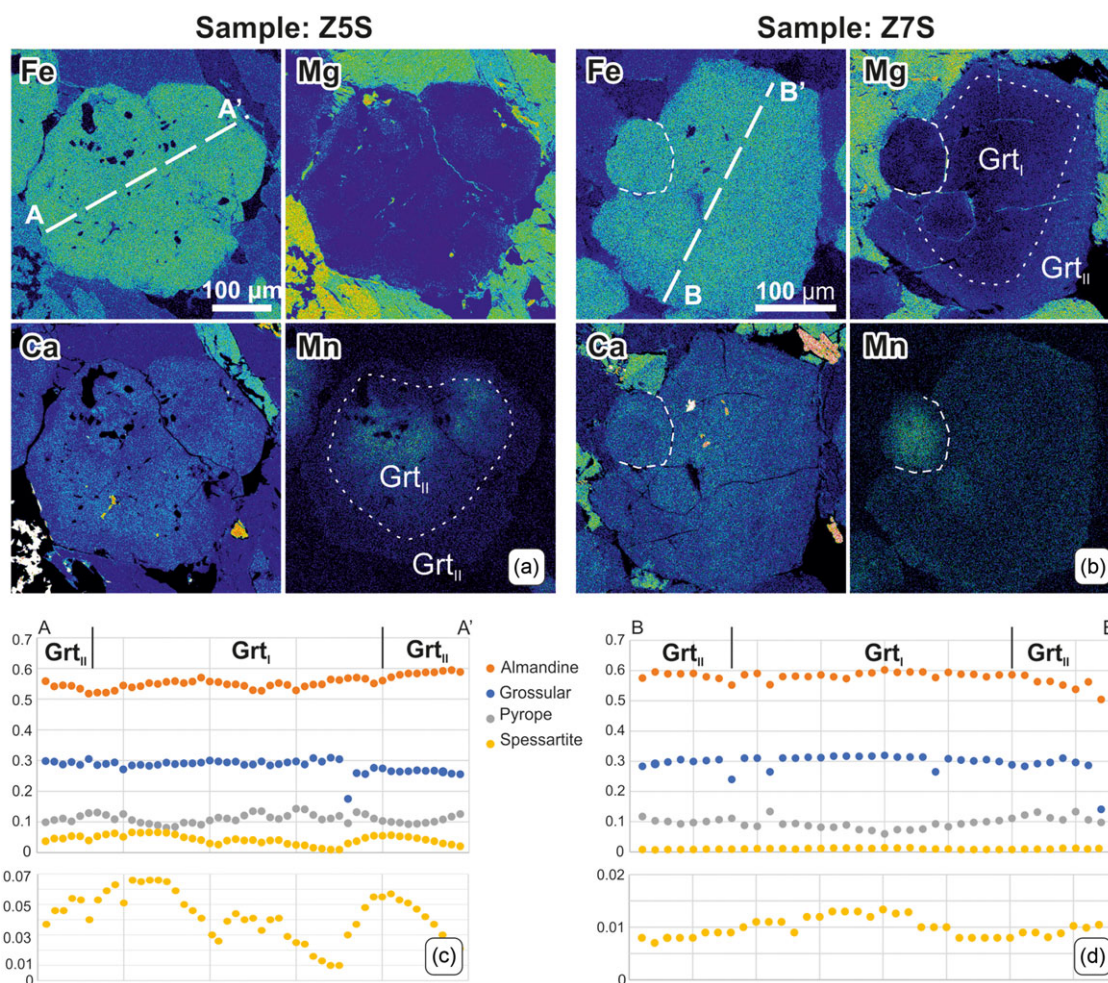


## 6. Mineral chemistry

Mineral phases representative of the crystallization/deformation stages reconstructed above were analysed to define the P-T conditions during the whole tectonometamorphic evolution of the NSMC.

X-rays elemental maps of garnet porphyroblasts (Figure 6) show a heterogeneous compositional variation. As shown in Figures 6c and 6d, the microstructural zoning characterized by inclusions-rich cores and clear rims reflects two crystallization events ( $\text{Grt}_I$  and  $\text{Grt}_{II}$ ). Both  $\text{Grt}_I$  and  $\text{Grt}_{II}$  show a poorly developed chemical zonation, with only a very slight increase of pyrope from cores to rims (Figure 6b). Grossular and spessartine are grossly homogeneous (exception done for microinclusions of the pre-eclogitic stage revealed by some spots). At a closer view, spessartine shows a patchy variation possibly derived from dissolution-precipitation phenomena associated with the occurrence of fluids (e.g. Giuntoli *et al.* 2018; Hyppolito *et al.* 2019). Garnet porphyroblasts are frequently fractured, as well observable in the Ca and Mg maps of Figure 6, allowing the penetration of fluids inside them.





**Figure 6.** (Colour online) WDS X-ray elemental maps of representative garnet porphyroblasts of the NSMC eclogites. Mn-rich cores are visible in both garnet porphyroblasts from samples Z5S (a) and Z7S (b). If Z7S garnet displays a continuous zoning, garnet from sample Z5S displays a patchy zoning, with irregular distribution of Ca and Mn abundances (a). The compositional profile obtained by means of quantitative WDS spot analyses along a line of Figure 6c related to the Mn-content clearly points to a polyphasic growth of garnet porphyroblasts. In the lower part of (c) and (d) the spessartine content is shown with an enlarged vertical scale to see the Sps variation along the profile.

The chemical composition of garnets from eclogites could be summarized as follows: Grt<sub>I</sub> (garnet cores) consists of Alm<sub>52-58</sub>Grs<sub>29-31</sub>Pyr<sub>9-15</sub>Sps<sub>1-7</sub>; Grt<sub>II</sub> (garnet rims) consists of Alm<sub>52-60</sub>Grs<sub>24-30</sub>Pyr<sub>10-13</sub>Sps<sub>0-3</sub> (representative EMPA of garnets are reported in Table 2).

Pre-eclogitic stage mineral phases are only preserved as inclusions in garnet cores (Grt<sub>I</sub>). Among them, phengitic white mica (Wm<sub>I</sub>) and amphibole (Amp<sub>I</sub>) are the most useful for the determination of P-T conditions during this stage. Wm<sub>I</sub> has a phengitic composition with Si contents ranging from 3.34 to 3.36 atoms per formula units (a.p.f.u.). Amphiboles included in garnet cores (Amp<sub>I</sub>) are sodic-calcic, with Si of 6.6-6.7 a.p.f.u. Following the classification scheme of Hawthorne & Oberti (2007), they could be classified as magnesio-katophorites. Other minerals of the same stage are epidote (Ep<sub>I</sub>), an almost pure pistacite (Table 2), quartz, ilmenite, and rare plagioclase (Pl<sub>I</sub>) displaying an albitic composition (Table 3). The observed microstructural relationships do not allow to assess if garnet was a stable phase during this stage (as proposed in Figure 5) or grew later, following increasing pressure conditions that lead to the eclogitic stage (Stage II, Figure 5).

Cpx<sub>II</sub> displays composition with a Na content expressed as jadeite from 0.36 to 0.46 mol% (Table 3), with increasing Na from core to rim (Figure 7a), resembling the prograde zonation of garnet.

The shape preferred orientation of Cpx<sub>II</sub>, and Wm<sub>II</sub> defines the eclogitic foliation. Wm<sub>II</sub> has higher Si content with respect to Wm<sub>I</sub> (Figure 7b), suggesting higher pressure conditions. The Si content of Wm<sub>II</sub> ranges from 3.36 to 3.40 a.p.f.u., approaching the celadonic end member. Rare Amp<sub>II</sub> crystals in eclogite, stable with the other HP mineral phases, also display compositions in agreement with the magnesio-katophorite compositional field (Figure 7d) like Amp<sub>I</sub>, but with slightly lower Si contents and  $X_{Mg}$  [Mg/(Mg+Fe<sup>2+</sup>)] values.

Eclogites of the NSMC are partially to completely retrogressed under amphibolites-facies conditions. Samples displaying the most complete retrogression during this stage were selected for EMPA. The Amp<sub>III</sub> partially replaces Amp<sub>II</sub> corresponding to a Mg-hornblende (Figure 7c) and its composition significantly differs from previous amphibole generation, becoming more Ca-rich (Figure 7c), plotting within the magnesio-hornblende field. The Na content of Amp<sub>III</sub> is lower (0.42-0.45 a.p.f.u.) with respect to Amp<sub>II</sub>, suggesting that they likely formed already during the exhumation after the pressure peak. Amp<sub>II</sub>, growing both at the expense of Amp<sub>I</sub> and in the matrix, has lower Ca and Si (6.15-6.25 a.p.f.u.). Fe-rich epidote (Ep<sub>III</sub>) is the Al-rich phase in place of garnet, suggesting lower pressure conditions during this stage, i.e. epidote-amphibolite facies.

**Table 2.** Representative EMPA of Grt porphyroblasts of the NSMC eclogites. (I) Garnet cores, (II) garnet rims

Mineral	Grt <sub>I</sub> (core)						Grt <sub>II</sub> (rim)					
	Z5S	Z5S	Z5S	Z7S	Z7S	Z7S	Z5S	Z5S	Z5S	Z7S	Z7S	Z7S
SiO <sub>2</sub>	38.63	38.45	38.57	40.66	38.5	38.38	38.21	38.73	38.66	39.02	38.79	38.88
TiO <sub>2</sub>	0.17	0.16	0.18	0.11	0.18	0.13	0.16	0.07	0.15	0.2	0.24	0.18
Al <sub>2</sub> O <sub>3</sub>	21.33	21.46	21.6	19.04	21.18	21.2	21.47	21.44	21.23	21	20.85	20.81
Cr <sub>2</sub> O <sub>3</sub>	0	0	0	0.01	0.02	0	0	0.04	0.05	0.06	0	0
FeO†	23.9	23.95	23.91	27.88	26.93	27.25	26.26	27.12	26.78	25.25	25.89	26.13
MnO	2.42	2.68	2.82	0.65	0.57	0.63	0.38	0.32	0.35	2.47	2.54	2.35
MgO	3.38	3.18	2.77	1.55	1.88	1.87	3.01	2.63	2.58	2.61	2.54	2.48
CaO	10.43	10.57	10.64	11.67	11.3	11.4	10.56	10.48	10.75	9.91	9.65	9.51
Na <sub>2</sub> O	0.07	0.05	0.02	0.01	0	0	0.06	0.02	0	0	0	0
K <sub>2</sub> O	0.02	0	0	0	0	0.01	0	0	0	0	0.01	0
Total	100.35	100.49	100.51	101.58	100.56	100.87	100.34	100.85	100.54	100.52	100.51	100.35
Si	3.02	3.01	3.02	3.19	3.03	3.02	3.00	3.03	3.04	3.07	3.06	3.07
Ti	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.00	0.01	0.01	0.01	0.01
Al	1.97	1.98	1.99	1.76	1.97	1.96	1.98	1.98	1.97	1.95	1.94	1.94
Cr	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Fe <sup>2+</sup>	1.56	1.57	1.57	1.83	1.77	1.79	1.72	1.77	1.76	1.66	1.71	1.73
Mn	0.16	0.18	0.19	0.04	0.04	0.04	0.03	0.02	0.02	0.16	0.17	0.16
Mg	0.39	0.37	0.32	0.18	0.22	0.22	0.35	0.31	0.30	0.31	0.30	0.29
Ca	0.87	0.89	0.89	0.98	0.95	0.96	0.89	0.88	0.90	0.84	0.82	0.80
Na	0.01	0.01	0.00	0.00	0.00	0.00	0.01	0.00	0.00	0.00	0.00	0.00
Grossular	0.29	0.29	0.29	0.32	0.31	0.31	0.28	0.29	0.30	0.27	0.27	0.26
Almandine	0.52	0.52	0.53	0.60	0.59	0.60	0.58	0.60	0.59	0.56	0.57	0.58
Pyrope	0.13	0.12	0.11	0.06	0.07	0.07	0.12	0.10	0.10	0.10	0.10	0.10
Spessartite	0.05	0.06	0.06	0.01	0.01	0.01	0.01	0.01	0.01	0.06	0.06	0.05

†Total iron as FeO.

## 7. <sup>40</sup>Ar/<sup>39</sup>Ar geochronology

Two samples have been investigated for the determination of the geochronological evolution of the NSMC (Figure 8 and Table 4). White mica separates from an eclogite (sample Z7S) and amphibole (Amp<sub>III</sub>) from an amphibolite (Z3S) have been analysed for <sup>40</sup>Ar/<sup>39</sup>Ar dating in order to constrain the timing of the peak pressure and the subsequent retrogression at epidote-amphibolites-facies conditions.

White mica (Wm<sub>II</sub>) grows aligned along the eclogitic foliation, so its recrystallization age broadly corresponds to the HP peak stage experienced by the NSMC eclogites. Taking into consideration the flat part of the age spectrum corresponding to homogeneous Ca/K values calculated from the <sup>37</sup>Ar/<sup>39</sup>Ar ratio (Figure 8) and the amount of gas released in the flat part of the spectrum, which corresponds to c. 84% of the total <sup>39</sup>Ar released, the age of white mica in Z7S is 191.14 ± 0.50 Ma (Figure 8a, b).

Large amphibole crystals belonging to the Amp<sub>III</sub> generation in completely retrogressed eclogites (i.e. amphibolite) have been selected for dating. The Ar release pattern of the Z3S sample (Figure 8c) is quite complex. The staircase pattern is likely related to the mixing trend of two amphibole generations, Amp<sub>II</sub> and Amp<sub>III</sub>, the latter partially overgrown at the expenses of Amp<sub>II</sub>

(see above), showing different Ca/K ratios (Figure 8d). Considering the chemical composition derived from the <sup>37</sup>Ar/<sup>38</sup>Ar, the cluster showing lower Ca/K ratios is referred to be 143.96 ± 5.82 Ma, whereas the cluster with higher Ca/K ratios has 194.38 ± 4.30 Ma. This age distribution accounts for the stability of amphibole during peak pressure condition and subsequent recrystallization with decreasing Ca/K ratio during retrogression. EMPA data indicate that Amp<sub>III</sub> has significantly lower Ca/K values with respect to Amp<sub>II</sub> (Table 3).

## 8. Thermodynamic modelling and metamorphic evolution

The microstructural analyses and the chemical composition of HP minerals in the studied (not retrogressed) eclogites indicate a prograde metamorphism mainly recorded by two stages of garnet crystallization (core, Grt<sub>I</sub>, and rim, Grt<sub>II</sub>), and a subsequent chemical zoning observed both in garnet rims (Grt<sub>II</sub>) and in omphacite (Cpx<sub>II</sub>). Moreover, the patchy garnet zoning (Figure 6c) was likely due to syn-deformation dissolution and reprecipitation processes mediated by metamorphic fluids. This is evidence of a complex metamorphic history that is difficult to reconstruct in detail from a thermodynamic modelling point of view.

**Table 3.** Representative EMPA of Cpx, Wm, Amp, Ep, and Pl of the NSCM eclogites. (I) pre-eclogite, (II) eclogite and (III) amphibolite metamorphic evolution stages

Mineral	Cpx <sub>I</sub>								Cpx <sub>II</sub>				Wm <sub>I</sub>					Wm <sub>II</sub>				
	Z5S	Z5S	Z5S	Z5S	Z5S	Z5S	Z5S	Z5S	Z3S	Z3S	Z5S	Z5S	Z5S	Z5S	Z5S	Z3S	Z3S	Z3S	Z5S	Z5S	Z5S	Z5S
SiO <sub>2</sub>	56.63	56.71	56.83	56.60	56.83	56.98	56.20	53.61	53.45	50.79	51.08	50.81	50.91	50.82	51.42	52.36	52.37	53.09	52.54	52.16	52.28	
TiO <sub>2</sub>	0.12	0.09	0.09	0.12	0.07	0.09	0.01	0.17	b.d.l.	0.53	0.53	0.50	0.52	0.55	0.54	0.61	0.61	0.59	0.49	0.56	0.57	
Al <sub>2</sub> O <sub>3</sub>	10.46	8.82	10.08	9.39	10.79	10.00	9.12	0.87	0.99	29.66	29.03	29.34	29.12	29.56	28.89	29.55	29.95	29.77	29.42	29.24	29.66	
Cr <sub>2</sub> O <sub>3</sub>	0.03	0.04	0.04	0.04	0.03	0.06	b.d.l.	b.d.l.	b.d.l.	b.d.l.	0.07	0.01	0.02	0.01	0.02	0.03	b.d.l.	0.04	b.d.l.	0.01	0.03	
FeO†	5.13	5.55	5.61	4.12	4.21	4.33	4.99	8.31	9.16	2.07	2.37	2.14	2.35	2.24	1.81	1.66	1.59	1.66	1.59	1.77	1.71	
MnO	0.00	0.02	0.03	0.01	0.00	0.05	0.02	0.24	0.11	0.01	0.07	0.07	0.07	0.03	0.05	b.d.l.	0.02	b.d.l.	b.d.l.	b.d.l.	0.01	
MgO	8.19	9.01	7.95	9.29	7.87	8.47	9.09	13.04	12.06	2.81	2.87	2.84	2.87	2.86	3.05	3.12	3.10	2.92	3.05	3.02	3.11	
CaO	13.79	15.04	14.08	15.47	13.59	14.43	15.09	23.58	23.69	0.02	0.02	0.02	0.01	0.01	b.d.l.	0.02	0.01	0.02	b.d.l.	0.01	0.03	
Na <sub>2</sub> O	6.81	5.74	6.66	5.51	6.64	6.38	5.73	0.31	0.59	0.89	0.84	0.81	0.82	0.84	0.81	0.69	0.75	0.72	0.68	0.69	0.68	
K <sub>2</sub> O	0.01	b.d.l.	b.d.l.	b.d.l.	b.d.l.	b.d.l.	0.01	0.01	b.d.l.	8.85	8.86	8.89	8.99	8.89	9.08	9.02	9.46	9.09	9.33	8.88	9.23	
Total	101.17	101.02	101.37	100.55	100.03	100.79	100.26	100.14	100.05	95.63	95.74	95.43	95.68	95.81	95.67	97.06	97.86	97.90	97.10	96.34	97.31	
Si	1.99	2.01	2.00	2.01	2.01	2.01	2.00	2.00	2.00	3.31	3.33	3.32	3.32	3.31	3.35	3.35	3.34	3.37	3.37	3.36	3.35	
Ti	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.01	–	0.03	0.03	0.02	0.03	0.03	0.03	0.03	0.03	0.03	0.03	0.02	0.03	
Al	0.43	0.37	0.42	0.39	0.45	0.42	0.38	0.04	0.05	2.28	2.23	2.26	2.24	2.27	2.22	2.23	2.25	2.23	2.23	2.22	2.24	
Cr	0.00	0.00	0.00	0.00	0.00	0.00	–	–	–	–	0.00	0.00	0.00	0.00	0.00	0.00	–	–	–	–	0.00	
Fe <sup>3+</sup>	0.04	0.03	0.04	0.00	0.01	0.02	0.01	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	
Fe <sup>2+</sup>	0.11	0.13	0.13	0.12	0.12	0.10	0.13	0.26	0.29	0.11	0.13	0.12	0.13	0.12	0.10	0.09	0.08	0.09	0.09	0.10	0.09	
Mn	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.01	0.00	0.00	0.00	0.00	0.00	0.00	0.00	–	0.00	–	–	–	0.00	
Mg	0.43	0.48	0.42	0.49	0.42	0.45	0.48	0.72	0.67	0.27	0.28	0.28	0.28	0.28	0.30	0.30	0.29	0.28	0.29	0.29	0.30	
Ca	0.52	0.57	0.53	0.59	0.52	0.55	0.58	0.94	0.95	0.00	0.00	0.00	0.00	0.00	–	0.00	0.00	0.00	0.00	0.00	0.00	
Na	0.47	0.39	0.45	0.38	0.46	0.44	0.40	0.02	0.04	0.11	0.11	0.10	0.10	0.11	0.10	0.09	0.09	0.09	0.08	0.09	0.08	
K	0.00	–	–	–	–	–	–	–	–	0.74	0.74	0.74	0.75	0.74	0.76	0.74	0.77	0.74	0.76	0.73	0.75	
Cation Sum	4.00	3.99	3.99	3.98	3.98	3.99	4.00	3.99	4.01	6.85	6.84	6.85	6.85	6.84	6.86	6.82	6.86	6.83	6.85	6.82	6.84	
X <sub>Mg</sub> =(Mg/Mg+Fe <sup>2+</sup> )	0.79	0.78	0.77	0.80	0.78	0.81	0.78	0.74	0.70	0.71	0.68	0.70	0.69	0.70	0.75	0.77	0.78	0.76	0.77	0.75	0.76	
Al <sup>(IV)</sup>	0.01	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	1.59	1.56	1.58	1.56	1.57	1.57	1.58	1.59	1.61	1.60	1.59	1.58	
Al <sup>(VI)</sup>	0.43	0.37	0.42	0.39	0.45	0.42	0.38	0.04	0.05	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	

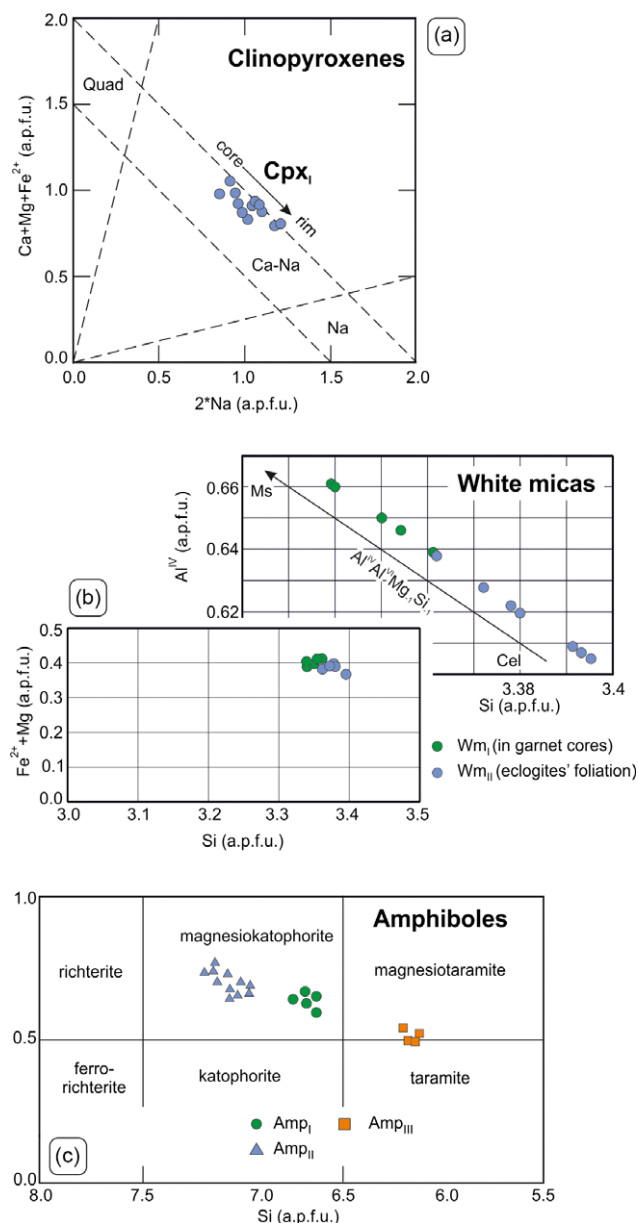
Mineral	Amp <sub>I</sub>			Amp <sub>II</sub>					Amp <sub>III</sub>					Ep <sub>I</sub>		Pl <sub>I</sub>		Pl <sub>III</sub>	
	Z3S	Z5S	Z5S	Z3S	Z3S	Z3S	Z5S	Z5S	Z3S	Z3S	Z3S	Z3S	Z3S	Z3S	Z7S	Z7S	Z5S	Z3S	Z5S
SiO <sub>2</sub>	46.37	45.98	46.44	51.21	50.36	49.22	49.02	49.93	47.54	42.14	47.63	39.45	39.15	39.72	39.42	68.81	68.66	60.91	
TiO <sub>2</sub>	0.42	0.38	0.43	0.36	0.39	0.37	0.46	0.72	0.38	0.28	0.38	0.18	0.09	0.09	0.03	b.d.l.	b.d.l.	b.d.l.	
Al <sub>2</sub> O <sub>3</sub>	14.68	14.69	14.59	11.69	11.96	11.86	12.26	11.78	11.86	19.10	12.15	29.56	28.10	29.37	28.55	19.63	19.67	25.08	
Cr <sub>2</sub> O <sub>3</sub>	b.d.l.	b.d.l.	b.d.l.	0.01	0.06	0.02	b.d.l.	0.05	0.11	0.04	0.05	b.d.l.	0.03	b.d.l.	0.01	0.03	0.00	0.02	
FeO†	12.10	12.11	12.01	8.86	9.26	11.23	11.18	10.54	12.63	15.00	12.63	6.11	7.50	6.11	7.39	0.27	0.29	0.41	
MnO	0.11	0.12	0.10	0.11	0.14	0.11	0.05	0.05	0.25	0.23	0.20	0.25	0.19	0.11	0.26	0.00	0.07	0.00	

(Continued)

**Table 3.** (Continued)

Mineral Sample	Amp <sub>I</sub>			Amp <sub>II</sub>					Amp <sub>III</sub>			Ep <sub>I</sub>				Pl <sub>I</sub>		Pl <sub>III</sub>
	Z3S	Z5S	Z5S	Z3S	Z3S	Z3S	Z5S	Z5S	Z3S	Z3S	Z3S	Z3S	Z3S	Z7S	Z7S	Z5S	Z3S	Z5S
MgO	11.37	11.31	11.38	14.18	13.81	12.34	12.60	12.66	12.14	8.70	12.47	0.07	0.06	0.08	0.00	0.01	b.d.l.	b.d.l.
CaO	9.42	9.54	9.45	8.85	9.07	9.05	9.05	9.06	11.53	11.05	11.90	23.49	23.17	23.50	23.02	0.31	0.36	5.90
Na <sub>2</sub> O	3.41	3.32	3.45	2.96	3.02	3.57	3.61	3.34	1.58	2.04	1.61	b.d.l.	b.d.l.	b.d.l.	0.01	10.99	10.92	7.87
K <sub>2</sub> O	0.48	0.43	0.45	0.35	0.38	0.44	0.42	0.38	0.21	0.45	0.19	b.d.l.	b.d.l.	b.d.l.	0.01	0.07	0.10	0.06
Total	98.36	97.88	98.30	98.58	98.45	98.21	98.65	98.51	98.23	99.03	99.21	99.11	98.29	98.98	98.70	100.12	100.07	100.25
Si	6.61	6.59	6.63	7.09	7.02	7.00	6.93	7.05	6.81	6.05	6.76	2.99	3.01	3.02	3.01	3.02	3.02	2.71
Ti	0.04	0.04	0.05	0.04	0.04	0.04	0.05	0.08	0.04	0.03	0.04	0.01	0.01	0.00	0.00	–	–	–
Al	2.47	2.48	2.45	1.91	1.97	1.99	2.04	1.96	2.00	3.23	2.03	2.64	2.54	2.63	2.57	1.02	1.02	1.31
Cr	–	–	–	0.00	0.01	0.00	–	0.01	0.01	0.01	0.01	–	0.00	–	0.00	0.00	0.00	0.00
Fe <sup>3+</sup>	0.30	0.32	0.26	0.36	0.31	0.10	0.20	0.07	0.27	0.57	0.26	0.39	0.48	0.39	0.47	0.00	0.00	0.00
Fe <sup>2+</sup>	1.14	1.13	1.17	0.67	0.77	1.24	1.12	1.18	1.24	1.23	1.24	0.00	0.00	0.00	0.00	0.01	0.01	0.02
Mn	0.01	0.01	0.01	0.01	0.02	0.01	0.01	0.01	0.03	0.03	0.02	0.01	0.01	0.01	0.02	0.00	0.00	0.00
Mg	2.42	2.42	2.42	2.93	2.87	2.62	2.65	2.66	2.59	1.86	2.64	0.01	0.01	0.01	0.00	0.00	–	–
Ca	1.44	1.47	1.45	1.31	1.36	1.38	1.37	1.37	1.77	1.70	1.81	1.91	1.91	1.91	1.88	0.01	0.02	0.28
Na	0.94	0.92	0.96	0.79	0.82	0.98	0.99	0.91	0.44	0.57	0.44	–	–	–	0.00	0.94	0.93	0.68
K	0.09	0.08	0.08	0.06	0.07	0.08	0.07	0.07	0.04	0.08	0.03	–	–	–	0.00	0.00	0.01	0.00
Cation Sum	15.47	15.47	15.48	15.17	15.24	15.44	15.43	15.35	15.25	15.35	15.29	7.97	7.97	7.97	7.96	5.00	5.00	5.00
X <sub>Mg</sub> =(Mg/Mg+Fe <sup>2+</sup> )	0.68	0.68	0.67	0.81	0.79	0.68	0.70	0.69	0.68	0.60	0.68	0.65	0.55	0.63	0.57	0.02	0.02	0.29
Al <sup>(IV)</sup>	1.39	1.41	1.37	0.91	0.98	1.00	1.07	0.95	1.19	1.95	1.24	0.35	0.45	0.37	0.43	0.98	0.98	0.70
Al <sup>(VI)</sup>	1.08	1.08	1.09	1.00	0.99	0.99	0.97	1.01	0.81	1.28	0.79							

†Total iron as FeO; b.d.l.: below detection limit.



**Figure 7.** (Colour online) (a) Clinopyroxene (Cpx<sub>II</sub>) composition, rims are enriched in Na. (b) The Si content of phengitic white micas increases from Wm<sub>I</sub>, preserved as inclusion in garnet cores to Wm<sub>II</sub> stable as HP phase. (c) Classification of Ca-Na amphibole composition in eclogites and amphibolites, see text for description.

Conventional geothermobarometric calculations have been performed on one of the best-preserved eclogites (Z5S) in order to compare the obtained estimates with the modelling results (e.g. Wei & Clarke, 2011). The occurrence of coexisting clinopyroxene and garnet stable in the peak metamorphic assemblage (Stage II) allows the use of the Fe-Mg exchange thermometer (Krogh Ravna, 1988, 2000) between the two phases. The occurrence of phengite also allows estimating the pressure using the geothermobarometer of Krogh Ravna & Terry (2004) developed for garnet + omphacite + phengite + kyanite + quartz in the KCMASH system. In the quartz stability field, the geothermobarometer consists of three independent reactions cross-cutting each other in a single invariant point (Figure 9a). One of the three reactions is kyanite and quartz-free ( $2 \text{ Grs} + \text{Pyr} + 3 \text{ Al-Cel} = 6 \text{ Di} + 3 \text{ Mus}$ ), therefore suitable for the kyanite-free eclogites of the NSCM. Minerals (garnet,

omphacite and phengite) in textural equilibrium showing no sign of retrogression have been selected and analysed with the EMP (Table 5), and the results are reported in Figure 9a and Table 5. The estimated pressure ranges from 1.90 to 2.15 GPa with T in the 510–600 °C using the Krogh Ravna (1988) thermometer. A restricted temperature interval, 510–540 °C, stems instead employing the Krogh Ravna (2000) version of the same thermometer.

With respect to conventional geothermobarometry, the reconstruction using forward modelling is in principle more robust even if the limitations of reliable solid solution models, particularly of amphibole, must be kept into consideration.

In order to check the role of CO<sub>2</sub> in the fluid and the stability of carbonates at the peak pressure as shown by the microstructural evidences, we constructed a T-X section of sample S5Z, based on the variation of X(CO<sub>2</sub>) and temperature, at P = 2 GPa (Figure 9b).

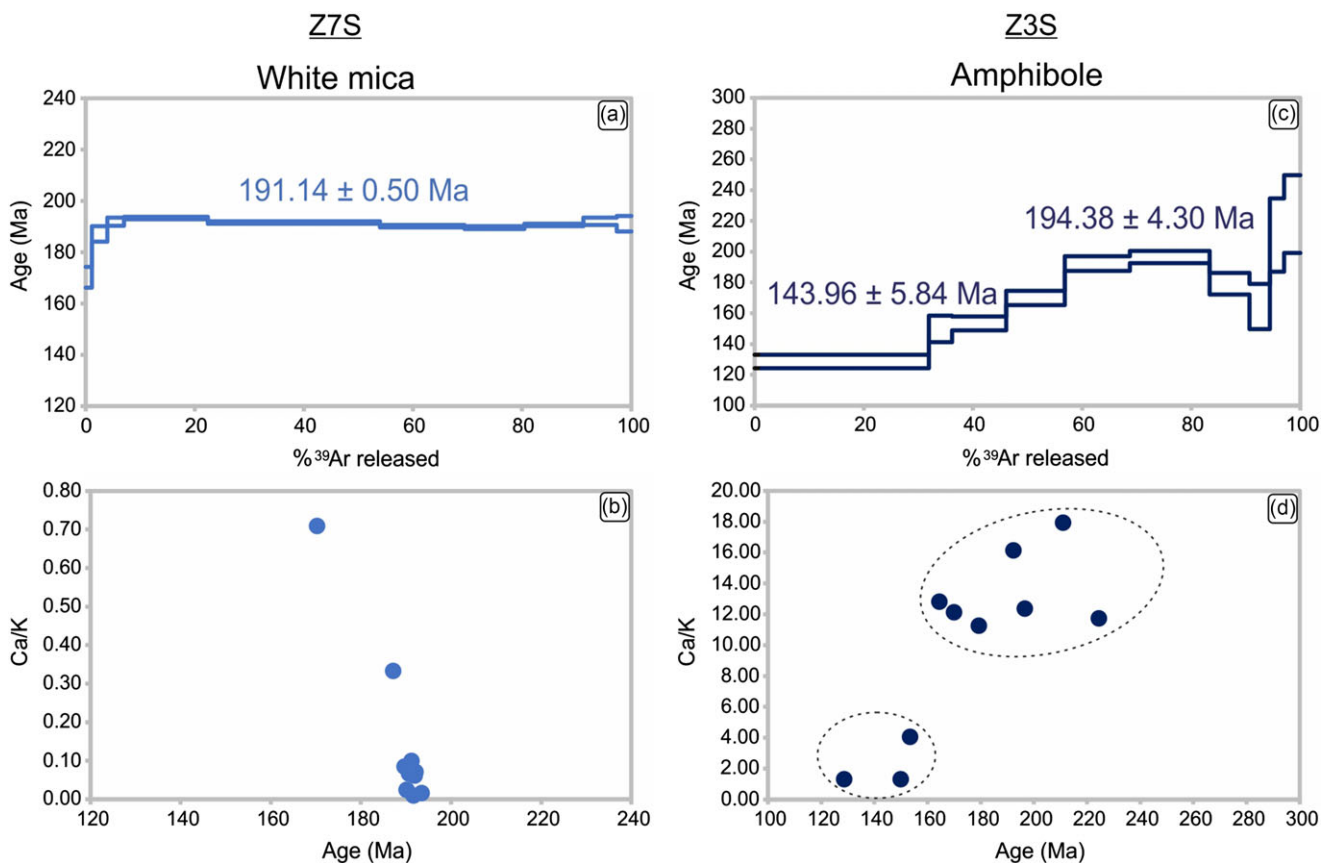
The amount of CO<sub>2</sub> strongly influences the stability of zoisite in a short range of temperature (550–650 °C, at X(CO<sub>2</sub>) up to 0.3). We therefore considered the phase assemblage garnet, omphacite, carbonate, quartz, zoisite, and rutile ( $\pm$  amphibole) as the possible mark of P-T range at X(CO<sub>2</sub>) up to 0.1, as demonstrated by microstructural evidences. The occurrence of amphibole strongly influences the temperature range of the mineral stability (550–600 °C for X(CO<sub>2</sub>) = 0.01–0.04 or 550–650 °C for X(CO<sub>2</sub>) = 0.015–0.03). If we assume that amphibole was in equilibrium at the peak and compare the garnet mol% compositions with those retrieved by the thermodynamic modelling (see properties in the Supplementary Material 1), we estimate that the peak P-T conditions recorded by our preserved eclogites range from 550 to 600 °C, at 2 GPa.

## 9. Discussion

### 9.a. P-T conditions of metamorphism

The studied samples show peak mineral assemblage and garnet composition compatible with pressures in the range between 1.8 and 2.1 GPa (Figure 9a), considering the conventional thermobarometry. More difficult is the determination of the equilibration temperature that ranges between 500 °C and 650 °C based on the occurrence of CO<sub>2</sub> in the fluid that may or may not stabilize carbonates among the HP mineral assemblages, along with the stability of amphibole at peak conditions. In addition, the comparison with experimental works on the phase stability of amphibole + zoisite in mafic assemblages (e.g. Schmidt & Poli, 1998; Forneris & Holloway, 2003) is in agreement with both the estimated temperatures, being stable between 500 and 700 °C. It must be noted that even if the bulk composition of our studied samples is slightly depleted in SiO<sub>2</sub> (46.71 vs 52.4 wt%) and enriched in MgO (8.30 vs 7.1 wt%) with respect to a MORB as reported in the experiments of Schmidt & Poli (1998), in the presence of fluids our phengite-bearing eclogites would have undergone partial melting at 1.8 GPa and 700 °C. The presence of fluids at peak conditions is demonstrated not only by the occurrence of hydrous phases but also by the microstructure of garnet zoning that resembles a coupled dissolution-precipitation and diffusional equilibration (Figure 6; Massonne & Li, 2020). However, the microstructures of our samples do not give evidence of such high temperatures and/or related symplectites growing during the retrogression, such as those encountered in the Central Italian Alps, at Borgo (Pellegrino *et al.* 2020).

The conditions of the formation of amphibole and clinozoisite-epidote porphyroblasts in eclogites have only rarely been addressed (Massonne, 2012 with references), but P-T conditions around



**Figure 8.** (Colour online)  $^{40}\text{Ar}/^{39}\text{Ar}$  age spectra and Ca/K vs. age diagrams for eclogite Z7S (a, b) and amphibolite Z3S (c, d). Dotted circles in (d) are referred to amphibole generations Amp<sub>II</sub> and Amp<sub>III</sub>.

2 GPa and 600 °C seem to be characteristic for their formation. These P estimates are similar to our data within the analytical error. Massonne (2012) suggested that the formation of amphibole and clinozoisite in eclogite is typical in subduction settings where deep burial of material from the overriding plate occurred, followed by fluid infiltration responsible for the formation of H<sub>2</sub>O-bearing HP mineral phases in former anhydrous phase assemblages. After the first hydration, the formation of hydrous porphyroblasts in HP rocks can continue also during their progressive exhumation, following resorption and compositional re-equilibration of existing porphyroblasts and the growth of new ones when the destabilization of hydrous phases occurs in response of P-T variations (e.g. Huang *et al.* 2020, 2023) in dynamic settings like subduction channels.

The eclogites from Dabie Shan in China (Massonne, 2012), Altiplano-Puna plateau in the central Andes (Allmendinger *et al.* 1997; Giese *et al.* 1999), Kaghan Valley in western Himalaya (Lombardo & Rolfo, 2000; Wilke *et al.* 2010), Cretaceous eclogites from the Austroalpine domain of the European Alps (e.g. Spalla *et al.* 1996; Thöni, 2006; Zanchetta *et al.* 2013b) may represent a geodynamic scenario similar to the one given by the eclogites of the North Shahrekord Metamorphic Complex. The clockwise P-T path (Figure 10d) constructed for the eclogite accounts for an initial pre-eclogite stage (stage I) preserved in garnet porphyroblasts, a HP eclogite facies metamorphic stage at 1.90–2.15 GPa and 550–600 °C and a subsequent epidote-amphibolite-facies retrograde stage. The estimated peak conditions fall for temperature in the range (470–630°C) proposed by Davoudian *et al.* (2008) and

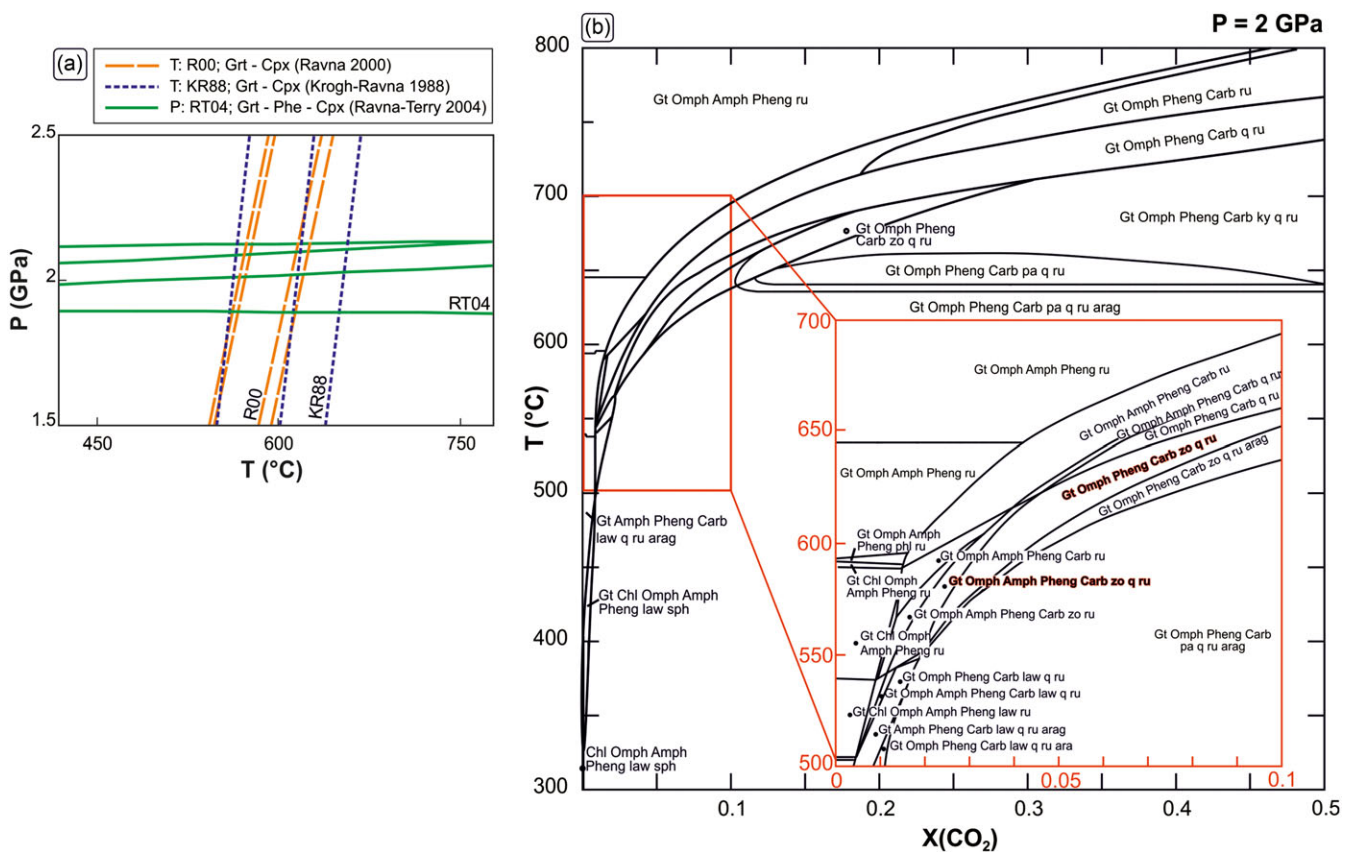
Davoudian *et al.* (2016) for the same NSMC eclogites. Our estimated pressures are instead lower by about 0.4–0.6 GPa, as previously calculated pressures are high as 2.5 GPa (Davoudian *et al.* 2008; Davoudian *et al.* 2016). We suggest that this higher pressures should be considered only as a maximum as a) the used chemical composition of HP phases was selected to maximize the Na content in omphacite, the Si content in phengite, and the Pyr and Grs activity in garnet; b) the equilibrium phase assemblage on which the used geothermobarometer (Krogh Ravna & Terry, 2004) considers the occurrence of kyanite, which is not present in the NSMC eclogites; c) the occurrence of carbonates (calcite and dolomite, Davoudian *et al.* 2016) has not been considered in the geothermobarometric calculations. Regarding these last points, we would stress how the occurrence of a fluid phase not formed only by pure H<sub>2</sub>O could significantly change the equilibrium phase assemblages as well as the composition of single minerals, as discussed before and shown in Figure 9b.

### 9.b. Timing and tectonic implications

The eclogites of the NSMC in the SSZ are the only record of Jurassic HP metamorphism identified in the Zagros orogen *sensu lato* (Davoudian *et al.* 2016), therefore being key rocks to the understanding of subduction-related processes that occurred in this area. The first question is if the  $^{40}\text{Ar}/^{39}\text{Ar}$  is a sound method to date the metamorphic peak in HP assemblage or if inherited Ar and fluid-aided retrogression could affect the Ar retention or loss. Besides the fact that many workers endeavoured to date HP and

**Table 4.**  $^{40}\text{Ar}/^{39}\text{Ar}$  data. All Ar isotope concentrations are given as mL. K, Ca and Cl concentrations are calculated from the sample mass and the  $^{39}\text{Ar}$ ,  $^{37}\text{Ar}$  and  $^{38}\text{Ar}$  concentrations, respectively.  $^{40}\text{Ar}^*$ =radiogenic  $^{40}\text{Ar}$ . All errors are  $1\sigma$

Sample	Step #	$^{40}\text{Ar}$ Tot	Err, $^{40}\text{Ar}$	$^{40}\text{Ar}^*$	Err, $^{40}\text{Ar}^*$	$^{39}\text{Ar}$	Err, $^{39}\text{Ar}$	% $^{39}\text{Ar}$	$^{38}\text{Ar}$	Err, $^{38}\text{Ar}$	$^{38}\text{ArCl}$	Err, $^{38}\text{Cl}$	$^{37}\text{Ar}$	Err, $^{37}\text{Ar}$	$^{36}\text{Ar}$	Err, $^{36}\text{Ar}$	Age	$1\sigma$ Err	Ca/K	Error Ca/K	Cl/K	Error Cl/K
Z7S	1	1.79E-08	4.50E-12	4.22E-09	2.65E-11	4.11E-11	4.20E-13	1.17E+00	1.37E-11	1.90E-14	4.69E-12	4.48E-14	1.50E-11	5.67E-14	4.57E-11	8.74E-14	170.27	2.04	0.70925	0.00772	0.02029	0.00028
J=0.000965	2	5.19E-08	2.10E-11	1.13E-08	7.92E-11	9.95E-11	3.92E-13	2.83E+00	2.95E-11	3.53E-14	2.80E-12	6.03E-14	1.71E-11	5.87E-14	1.36E-10	2.56E-13	187.17	1.51	0.33293	0.00174	0.00500	0.00011
m=0.00616 g	3	1.67E-08	3.60E-12	1.25E-08	9.60E-12	1.07E-10	4.23E-13	3.05E+00	4.32E-12	1.02E-14	5.28E-13	7.14E-14	3.38E-12	4.96E-14	1.41E-11	2.98E-14	191.92	0.77	0.06126	0.00093	0.00088	0.00012
	4	6.82E-08	3.60E-11	6.33E-08	3.75E-11	5.39E-10	5.66E-13	1.54E+01	1.04E-11	1.78E-14	1.54E-12	5.25E-14	4.33E-12	5.00E-14	1.67E-11	3.52E-14	193.45	0.24	0.01558	0.00018	0.00051	0.00002
	5	1.33E-07	9.90E-11	1.28E-07	9.96E-11	1.11E-09	1.25E-12	3.15E+01	1.74E-11	2.92E-14	2.58E-12	5.31E-14	5.21E-12	5.07E-14	1.66E-11	3.65E-14	191.64	0.26	0.00913	0.00009	0.00042	0.00001
	6	6.86E-08	3.30E-11	6.27E-08	3.53E-11	5.44E-10	5.67E-13	1.55E+01	1.09E-11	1.91E-14	1.49E-12	5.37E-14	6.99E-12	5.11E-14	1.95E-11	4.21E-14	190.19	0.23	0.02491	0.00018	0.00049	0.00002
	7	5.10E-08	1.71E-11	4.44E-08	2.18E-11	3.86E-10	5.11E-13	1.10E+01	9.45E-12	1.64E-14	1.22E-12	5.30E-14	1.69E-11	5.81E-14	2.21E-11	4.50E-14	189.69	0.27	0.08474	0.00031	0.00056	0.00002
	8	5.56E-08	2.31E-11	4.37E-08	3.27E-11	3.78E-10	4.26E-13	1.08E+01	1.27E-11	2.03E-14	1.21E-12	5.03E-14	1.31E-11	5.60E-14	3.99E-11	7.76E-14	190.70	0.26	0.06715	0.00030	0.00057	0.00002
	9	5.06E-08	5.40E-11	2.51E-08	7.44E-11	2.16E-10	4.60E-13	6.15E+00	1.92E-11	3.12E-14	8.90E-13	5.87E-14	7.92E-12	5.23E-14	8.53E-11	1.71E-13	192.14	0.70	0.07119	0.00049	0.00073	0.00005
	10	4.80E-08	2.43E-11	1.08E-08	7.43E-11	9.31E-11	3.62E-13	2.65E+00	2.49E-11	3.17E-14	4.29E-13	5.87E-14	4.77E-12	5.08E-14	1.25E-10	2.35E-13	191.23	1.51	0.09933	0.00113	0.00082	0.00011
Conc (g/g)	K	Cl	Ca																			
		8.45E-02	7.44E-05	4.42E-03																		
Z3S	1	4.33E-08	2.16E-11	6.59E-09	1.09E-10	8.61E-11	3.63E-13	3.19E+01	4.19E-11	6.76E-14	1.78E-11	8.30E-14	5.87E-11	2.06E-13	1.23E-10	3.57E-13	128.60	2.19	1.32158	0.00725	0.03670	0.00023
J=0.000965	2	1.58E-09	5.40E-13	1.03E-09	2.96E-12	1.14E-11	3.28E-13	4.24E+00	8.55E-13	5.30E-15	3.85E-13	1.86E-13	7.74E-12	1.42E-13	1.85E-12	9.76E-15	149.84	4.32	1.31316	0.04469	0.00599	0.00289
m=0.00799 g	3	4.33E-09	8.70E-13	2.47E-09	6.29E-12	2.69E-11	3.88E-13	9.97E+00	3.78E-12	9.92E-15	2.31E-12	7.88E-14	5.63E-11	1.97E-13	6.24E-12	2.09E-14	153.44	2.25	4.06396	0.06044	0.01530	0.00057
	4	4.52E-09	9.30E-13	2.94E-09	5.77E-12	2.88E-11	3.88E-13	1.07E+01	5.46E-12	1.30E-14	4.11E-12	7.19E-14	1.80E-10	4.31E-13	5.34E-12	1.91E-14	169.96	2.32	12.13909	0.16667	0.02535	0.00056
	5	4.87E-09	1.05E-12	3.74E-09	4.46E-12	3.22E-11	3.89E-13	1.19E+01	6.32E-12	1.43E-14	5.20E-12	6.80E-14	2.67E-10	6.06E-13	3.86E-12	1.45E-14	192.25	2.34	16.13559	0.19898	0.02871	0.00051
	6	5.99E-09	1.29E-12	4.68E-09	5.24E-12	3.94E-11	3.89E-13	1.46E+01	7.42E-12	1.51E-14	6.11E-12	6.13E-14	2.50E-10	5.71E-13	4.46E-12	1.70E-14	196.50	1.96	12.37108	0.12574	0.02759	0.00039
	7	3.31E-09	7.50E-13	2.13E-09	4.67E-12	1.98E-11	3.88E-13	7.32E+00	3.48E-12	9.95E-15	2.49E-12	8.60E-14	1.14E-10	2.99E-13	3.99E-12	1.54E-14	179.22	3.56	11.25716	0.22395	0.02246	0.00089
	8	2.40E-09	5.70E-13	9.95E-10	5.11E-12	1.01E-11	4.48E-13	3.74E+00	2.39E-12	7.63E-15	1.38E-12	9.62E-14	6.66E-11	2.17E-13	4.72E-12	1.70E-14	164.36	7.36	12.83210	0.57222	0.02422	0.00201
	9	2.78E-09	6.60E-13	8.90E-10	6.44E-12	6.97E-12	3.88E-13	2.58E+00	2.64E-12	8.59E-15	1.35E-12	9.79E-14	6.41E-11	2.10E-13	6.36E-12	2.14E-14	210.93	11.91	17.94409	1.00647	0.03450	0.00316
	10	9.47E-09	1.98E-12	1.11E-09	2.47E-11	8.13E-12	4.18E-13	3.01E+00	6.85E-12	1.50E-14	1.47E-12	6.77E-14	4.90E-11	1.85E-13	2.80E-11	8.24E-14	224.39	12.61	11.73584	0.60729	0.03222	0.00223
Conc (g/g)	K	Cl	Ca																			
		4.99E-03	1.41E-04	4.01E-02																		



**Figure 9.** (Colour online) (a) Results of P-T estimate using conventional geothermobarometry with the composition of mineral pairs reported in Supplementary Material 2 (b) T-X chemical sections of the eclogite Z5S (see composition in the ‘Methods’ section), where  $X(\text{CO}_2) = \text{CO}_2/(\text{CO}_2 + \text{H}_2\text{O})$ . The red box is a zoom of the phase assemblages in the range of 500 – 700 °C and  $X(\text{CO}_2)$  between 0 and 0.1. Abbreviations for minerals and compositional end-members after Connolly (2005).

UHP metamorphism through  $^{40}\text{Ar}/^{39}\text{Ar}$  on phengite and amphibole (Halama *et al.* 2014 and references therein), a long discussion about inherited and excess Ar is reported in Villa *et al.* (2014), who demonstrated that excess Ar was never credibly documented supported also by the absence of a grain size effect and by multichronometry (e.g. Lu-Hf and Rb-Sr). Regarding the fluid-aided retrogression, since the formation of retrograde minerals involves a recrystallization, a complete loss of Ar occurs, even at very low temperature (Maineri *et al.* 2003). Taking into account all these aspects,  $^{40}\text{Ar}/^{39}\text{Ar}$  on phengite and amphibole is still a robust method for dating high-pressure rocks returning a minimum age of peak metamorphism.

Several contrasting models have been proposed to explain the formation and geodynamic evolution of the SSZ metamorphic belt (Hassanzadeh & Wernicke, 2016 and references therein; Shafaii Moghadam *et al.* 2020; Maghdour-Mashhour *et al.* 2021; Gharibnejad *et al.* 2023), mostly concerning the onset of the Neo-Tethys subduction event and the related magmatism without taking into account the occurrence of Lower Jurassic eclogites.

The MORB-like affinity of the NSMC eclogites (Davoudian *et al.* 2006) is testified also by data presented in this work (Table 1). However, no geochronological constraints exist on the protoliths’ age. As the NSMC eclogites, as well as other units forming the SSZ metamorphic belt, show a clear Gondwanan affinity (Fergusson *et al.* 2016), they should be considered to be originally in an upper plate position during the onset of the subduction of the Neo-Tethys. A mechanism capable to drag slices of continental crust

derived from the upper plate at HP conditions should then be invoked to explain the formation of the NSMC eclogites. In the early phases of evolution of an orogen, a primary role may be played by tectonic erosion during subduction. Tectonic erosion has been widely recognized in oceanic subductions (e.g. Von Huene & Scholl, 1991; Barth & Schneiderman, 1996; Clift & Vannucchi, 2004; Clift & Hartley, 2007; Pastor-Galán *et al.* 2021) and rarely documented in continental subduction zones (e.g. Yin *et al.* 2007; Lu *et al.* 2018) or in back-arc basins (Peng *et al.* 2022). Peng *et al.* (2022) reported the occurrence of retrograde eclogites, characterized by peak metamorphic mineral assemblages with garnet, omphacite, rutile and quartz, within the Amdo microcontinent in central Tibet. These authors suggest that tectonic erosion associated with the subduction of the Amdo microcontinent beneath the Tethys Ocean accounts for the deep subduction induced by tectonic erosion of the overriding plate. Lu *et al.* (2018) also reported evidence of tectonic erosion related to continental subduction in north-western China. In this scenario, the eclogites from NSMC can thus be interpreted as lithologically heterogeneous tectonic slices derived from the upper plate, eroded and deeply buried *via* subduction channel on top of the subducting slab (Figure 10a). They subsequently underwent eclogite facies metamorphism (Figure 10b), exhumed along an exhumation channel and (re-)coupled with the crust of the overriding plate experiencing amphibolite-facies metamorphism (Figure 10c).

An additional hypothesis, that is also coherent with the upper plate derivation of the NSMC eclogites, is the subduction of tectonic slices derived from the hyperextended continental margin



**Table 5.** Compilation of ages of Early-Middle Jurassic magmatic activity in the SSZ

Location	Rock type	Mineral	Method	Age (Ma)	error	Reference
NW sector						
Bolban Abad	Granite and diorite	Zrn	U-Pb	146	2	Zhang et al. 2018
Bolban Abad	Granite and diorite	Zrn	U-Pb	148	2	Zhang et al. 2018
Suffi Abad	Granite and diorite	Zrn	U-Pb	156	0.6	Mahmoudi et al. 2011
Kangareh-Taghiabad	Gabbro	Zrn	U-Pb	148	4	Azizi et al. 2015a
Kangareh-Taghiabad	Gabbro	Zrn	U-Pb	158	10	Azizi et al. 2015a
Ghalaylan	Granite	Zrn	U-Pb	156	6	Azizi et al. 2015b
Ghalaylan	Granite	Zrn	U-Pb	158	2	Azizi et al. 2015b
Ghalaylan	Gabbro and monzonite	Zrn	U-Pb	149	0.2	Mahmoudi et al. 2011
Ghalaylan	Gabbro and monzonite	Zrn	U-Pb	151	0.2	Mahmoudi et al. 2011
Ghorveh	Granite and gabbro	Zrn	U-Pb	160.6	4	Yajam et al. 2015
Ghorveh	Granite and gabbro	Zrn	U-Pb	141.3	2	Yajam et al. 2015
Ghorveh	Granite and diorite	Zrn-Ttn	U-Pb/U-Th-Pb	146	2	Zhang et al. 2018
Ghorveh	Granite and diorite	Zrn-Ttn	U-Pb/U-Th-Pb	151	1	Zhang et al. 2018
Ghorveh	Porphyritic granite and diorite	Zrn	U-Pb	165	0.2	Mahmoudi et al. 2011
Ghorveh	Porphyritic granite and diorite	Zrn	U-Pb	165	0.4	Mahmoudi et al. 2011
Gorveh	Granitoid	Zrn	U-Pb	156.5	0.6	Mahmoudi et al. 2011
Gorveh	Granitoid	Zrn	U-Pb	149.3	0.2	Mahmoudi et al. 2011
Alvand, Hamedan	Granitoid	Zrn	U-Pb	165.02	0.2	Mahmoudi et al. 2011
Alvand, Hamedan	Leucogranite and granite	Zrn	U-Pb	164	2	Chiu et al. 2013
Alvand, Hamedan	Leucogranite and granite	Zrn	U-Pb	165	2	Chiu et al. 2013
Alvand, Hamedan	Gabbro	Zrn	U-Pb	166.5	1.9	Shahbazi et al. 2010
Alvand, Hamedan	Granite	Zrn	U-Pb	163.9	161.7	Shahbazi et al. 2010
Alvand, Hamedan	Leucogranite	Zrn	U-Pb	154.4	1.3	Shahbazi et al. 2010
Malayer	Granite, tonalite, granodiorite, syenogranite	Zrn	U-Pb	162	0.7	Ahadnejad et al. 2011
Malayer	Granite, tonalite, granodiorite, syenogranite	Zrn	U-Pb	187	0.3	Ahadnejad et al. 2011
Boroujerd	Quartz diorite, granodiorite	Zrn	U-Pb	170	0.2	Ahmadi Khalaji et al. 2007
Boroujerd	Quartz diorite, granodiorite	Zrn	U-Pb	172	1.5	Ahmadi Khalaji et al. 2007
Boroujerd	Granitoid	Zrn	U-Pb	169	1	Mahmoudi et al. 2011
Boroujerd	Granitoid	Zrn	U-Pb	169	172	Ahmadi Khalaji et al. 2007
Nezamabad	Monzogranite	Zrn	U-Pb	170	0.3	Mahmoudi et al. 2011
Central sector						
Astaneh	Granite	Zrn	U-Pb	168	1	Mahmoudi et al. 2011
Astaneh	Granitoid	Zrn	U-Pb	167.5	1	Mahmoudi et al. 2011
Astaneh	Granite	Zrn	U-Pb	170.7	1	Ahmadi Khalaji et al. 2007
Darjune, Azna	Gabbro	Zrn	U-Pb	170	3	Shakerardakani et al. 2015
Aligoordaz	Granite	Zrn	U-Pb	165	5	Esna-Ashari et al. 2012
Aligoordaz	Boninite-like			169.8	0.64	Esna-Ashari et al. 2016

(Continued)

**Table 5.** (Continued)

Location	Rock type	Mineral	Method	Age (Ma)	error	Reference
Aligoordaz	Granite	Zrn	U-Pb	164	2.1	Chiu <i>et al.</i> 2013
Kolah Ghazi, S Esfahan	Granodiorite, monzogranite and syenogranite	Zrn	U-Pb	175.8	3	Bayati <i>et al.</i> 2017
Kolah Ghazi, S Esfahan	Granodiorite, monzogranite and syenogranite	Zrn	U-Pb	168	1	Bayati <i>et al.</i> 2017
Tootak, Bavanat	Aplite granite	Zrn	U-Pb	173	9	Hosseini <i>et al.</i> 2012
Zayanderud	Pegmatite	Zrn	U-Pb	176.5	1.65	Jamali Ashtiani <i>et al.</i> 2020
Quri Complex, Chah Dozdan, Chah Bazargan	Anorthosite, quartz diorite, gabbro, granite	Zrn	U-Pb	171	2	Fazlnia, 2007; Fazlnia <i>et al.</i> 2013
Quri Complex, Chah Dozdan, Chah Bazargan	Anorthosite, quartz diorite, gabbro, granite	Zrn	U-Pb	173	2	Fazlnia, 2007; Fazlnia <i>et al.</i> 2013
Quri Complex, Chah Dozdan, Chah Bazargan	Anorthosite, quartz diorite, gabbro, granite	Zrn	U-Pb	164	8	Fazlnia, 2007; Fazlnia <i>et al.</i> 2013
Chah Ghand	Granodiorite-gabbro	Amp	K-Ar	163	4	Sheikholeslami <i>et al.</i> 2008
Sargaz, NW Jiroft	Granite	Zrn	U-Pb	175.2	1.6	Chiu <i>et al.</i> 2013
SE sector						
Sargaz, NW Jiroft	Granite	Zrn	U-Pb	175	2	Chiu <i>et al.</i> 2013
Remeshk-Ramak	Granite, diorite, trondhjemite, plagiogranite	Zrn	U-Pb	170	1	Hunziker <i>et al.</i> 2015
Remeshk-Ramak	Granite, diorite, trondhjemite, plagiogranite	Zrn	U-Pb	176	1	Hunziker <i>et al.</i> 2015
Remeshk-Ramak	Granite, diorite, trondhjemite, plagiogranite	Zrn	U-Pb	153	1	Hunziker <i>et al.</i> 2015
Remeshk-Ramak	Granite, diorite, trondhjemite, plagiogranite	Zrn	U-Pb	166	1	Hunziker <i>et al.</i> 2015
Makran	Trondhjemite	Zrn	U-Pb	156.4	1.7	Hunziker <i>et al.</i> 2011
Makran	Trondhjemite	Zrn	U-Pb	160.5	1.4	Hunziker <i>et al.</i> 2011

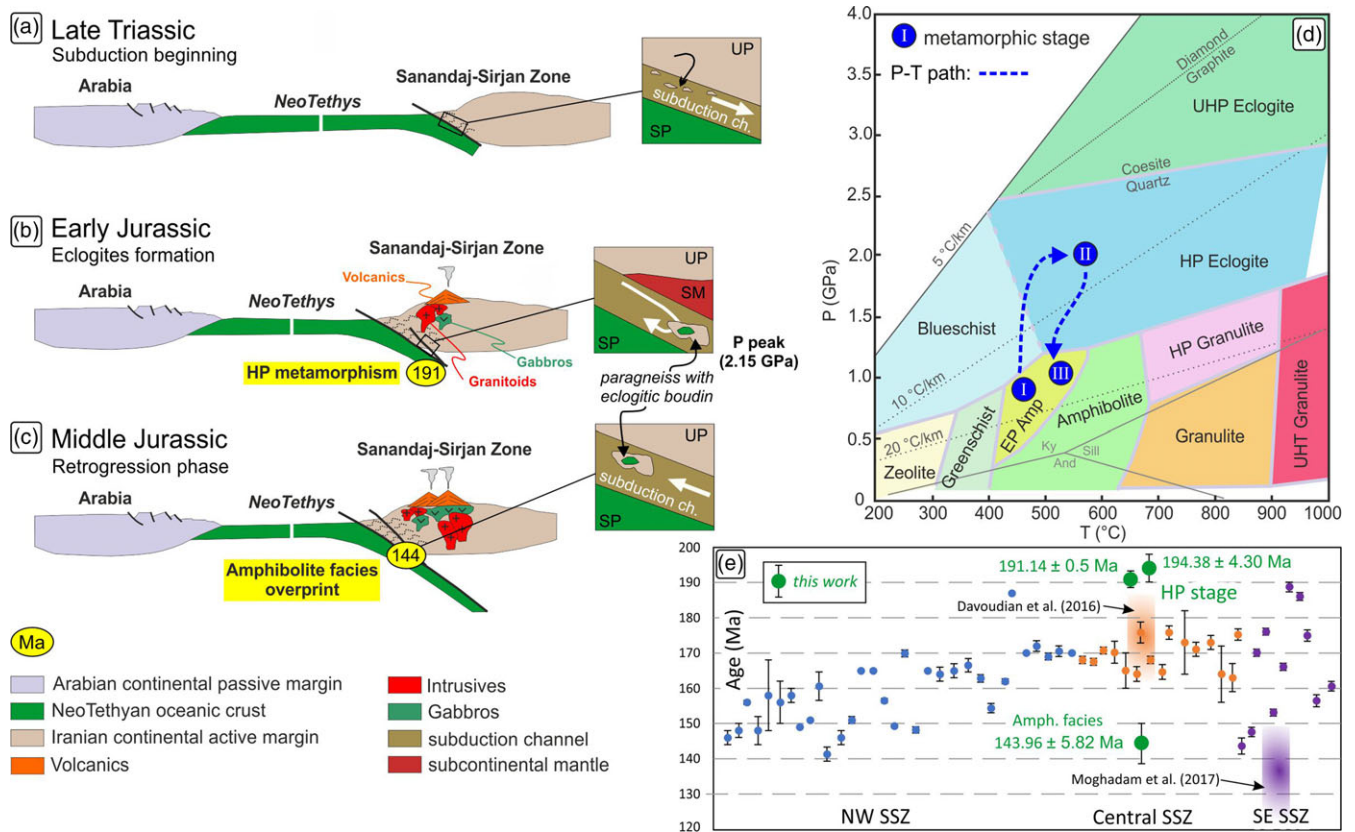
at the ocean-continent transition (Manatschal *et al.* 2022). However, in order to lean towards this hypothesis, remnants of (meta)sedimentary sequences and fossil structures typical of rifted margins (Manatschal & Chenin, 2023) should have been recognized.

The development of a volcanic arc magmatism (Figure 10a-c) of calc-alkaline affinity either in an island arc setting (i.e. Japan-like arc) or on a continental margin (i.e. Andean-type arc) has been broadly considered as an indirect evidence that subduction of the Neo-Tethys beneath Central Iran began in the Early Jurassic (Fergusson *et al.* 2016; Hassanzadeh & Wernicke, 2016) or even at the very end of the Triassic (Berberian & King, 1981). Indeed, in addition to ubiquitous Lower Jurassic intrusive bodies (187 Ma the oldest, with a climax around 170 Ma: Figure 10e and Table 5), diffuse coeval calc-alkaline volcanism and volcanoclastic sedimentation became prevalent in the SSZ, which is the key proof of the transition from passive margin sedimentation to the growth of a magmatic arc along an active margin (Hassanzadeh & Wernicke, 2016). The occurrence of eclogites in the SSZ supports the onset of an active convergent margin on the southwestern side of Central Iran, as was already supposed in the palaeogeographic reconstructions of Berberian & King (1981). They suggested that a subduction zone was active since the Late Triassic and related to

the consumption of what they called the ‘High-Zagros Alpine ocean’, dividing Central Iran from Arabia. The subduction margin developed in response to the closure of the Palaeo-Tethys to the north and the accretion of Central Iran to the southern Eurasia margin. Subduction moved from the Palaeo-Tethys suture, after the Eo-Cimmerian orogenic event (Zanchi *et al.* 2009a, 2009b; Zanchetta *et al.* 2013a), to the south. Evidence of the onset of an active margin along the SSZ are the andesitic-basaltic volcanism and granitoid intrusions exposed under the Jurassic and Cretaceous sediments of the SSZ (Berberian & King, 1981).

Available data (Figure 10e and Table 5) demonstrate that the magmatic history is continuous in the time span from c. 187 Ma (Ahadnejad *et al.* 2011) to c. 141 Ma (Yajam *et al.* 2015; Figure 10e), with some variations in geochemical affinity (see Maghdour-Mashhour *et al.* 2021 for an updated review).

The beginning of subduction of the Neo-Tethys started early in the whole SSZ, from the northern to the southern sectors (Sheikholeslami, 2015), probably starting in the Late Triassic (Figure 10a). Until now, the subduction of the Neo-Tethys beneath the SSZ was traced back up to ca. 185 Ma (Davoudian *et al.* 2016) and 210–192 Ma (Wan *et al.* 2023), based on the oldest eclogite ages of the NSMC in the central sector of the SSZ. The ages from Wan *et al.* (2023) closely match our ages, but have been obtained on



**Figure 10.** (Colour online) A schematic diagram showing the tectonic evolution of the SSZ from the (a) Late Triassic to the (b) Early and (c) Middle Jurassic. The white arrows in boxes (a, b, c) show the prevalent direction of materials in the subduction channel; the black arrow (a) points to the upper plate-derived rocks dragged in the subduction channel. (d) P-T path (in blue) for our studied eclogites derived from pseudosection modelling and conventional geothermobarometry. (e) Age distribution of magmatic activity in the SSZ. The orange box refers to the age of phengite in the eclogite of the NSMC from Davoudian *et al.* (2016); the violet box refers to the age of SE Zagros eclogite from Moghadam *et al.* (2017). UP: upper plate; SM: subcontinental mantle; SP: subducting plate.

rutile grains in which textural equilibrium with other syn-eclogite mineral phases (i.e. garnet and omphacite) is poorly constrained. Moreover, they consider the obtained lower intercept age as representative of peak metamorphism, even if each single age of different rutile grains displays  $1\sigma$  error always larger than 30 Ma. Phengite in our samples is instead clearly related to peak metamorphism as it is in equilibrium with garnet and omphacite, as testified by microstructural fabric, constraining a minimum age for peak metamorphism. Moreover, we constrain the forward part of the P-T path, corresponding to the onset of subduction.

Magmatism started slightly earlier in the SE SSZ (Figure 10e) around 188-189 Ma (Shafaii Moghadam *et al.* 2017). From SE to NW, the onset of subduction-related magmatism becomes younger (Figure 10e), starting only at 155-160 Ma in the NW sector of the SSZ.

Our new petrological and geochronological data testify that Neo-Tethys subduction was active at least since 191-194 Ma (Figures 8a and 10b), in the earliest Jurassic, just after the end of the Eo-Cimmerian orogeny. The oldest robust geochronological data on the onset of a calc-alkaline magmatic activity in the SSZ trace back to ca. 190 Ma (Figure 10). As the onset of volcanic arc magmatism in a supra-subduction setting requires several Myrs or tens of Myrs to flare up (e.g. Stein & Stein, 1992), we should consider 191 Ma as the minimum age of subduction initiation along the SSZ. P-T estimates of the pressure peak recorded by the NSMC eclogites presented in this study, together with the ones of Davoudian *et al.* (2008), suggest a geothermal gradient  $< 10^{\circ}\text{C}/\text{km}$ ,

pointing to an already mature subduction zone with a cold thermal regime (Cloos, 1993).

In the Middle Jurassic (Figure 10c), the retrogression phase was going on, developing amphibolites at the expense of the eclogites. This phase is constrained at 144 Ma by  $^{40}\text{Ar}/^{39}\text{Ar}$  age on amphibole (Figures 8c and 10b). The occurrence of Jurassic metamorphism in the SSZ preceding the Iran-Arabia collision is not limited to the NSMC. At its NW termination, the NSMC is in contact with the Hamadan Phyllite, a metamorphic complex mainly composed of phyllites and micaschists. The occurrence of polyphase deformation and of aluminosilicates such as andalusite, sillimanite and staurolite in several areas of the complex (Mohajjel *et al.* 2006) suggests that the Hamadan Phyllite underwent medium-grade metamorphism. Even if no direct dating is available, the intrusion ages of undeformed granitoids that cluster around 170 Ma (Ahmadi Khalaji *et al.* 2007; Mahmoudi *et al.* 2011) constrain the metamorphism in pre-Middle Jurassic times (Fergusson *et al.* 2016), so likely almost coeval with the HP event in the NSMC and subsequent amphibolite-facies retrogression.

Our new data imply that the Neo-Tethys subduction began quite before 191 Ma, therefore suggesting a subduction initiation at least 7 Myrs earlier compared to what was supposed previously (Davoudian *et al.* 2016). The age of HP eclogite facies metamorphism therefore preceded the onset of magmatism in the SSZ as should be expected for subduction-related magmatism.

Our results provide the first complete prograde-retrograde P-T-t path of eclogites in the SSZ (Figure 10d), constraining the

onset of eclogite facies metamorphism at 191 Ma (Figure 10b) and its duration up to subsequent amphibolite-facies retrogression at 144 Ma (Figure 10c). The subduction of the Neo-Tethys was accompanied by tectonic erosion from the Gondwanan overriding plate (Figure 10a), bringing the upper plate-related material to experience eclogite facies metamorphism (191 Ma, Figure 10b) and subsequent exhumation undergoing amphibolite-facies metamorphism (144 Ma, Figure 10c).

The Sinemurian age of the NSMC eclogites indicates that immediately after the end of the Eo-Cimmerian orogenesis, the subduction of the Neo-Tethys began along the southern border of the Iranian blocks, represented by the SSZ (Fergusson *et al.* 2016). The NSMC eclogites likely derived from crustal fragment of the upper plate (SSZ) that were scraped off by tectonic erosion by the subducting Neo-Tethys oceanic slab.

Evidence of a post-Cimmerian compression and plate reorganization in the Iranian geodynamic scenario is widespread. A 'subduction jump' from the Palaeo-Tethys margin to the Neo-Tethys one, i.e. the SSZ, could explain compressional deformation recorded in several parts of Central Iran in the Early to Middle Jurassic (Bagheri & Stampfli, 2008; Rahmati-Ilkhchi *et al.* 2011).

## 10. Conclusions

1. The eclogites of the North Shahrekord Metamorphic Complex (NSMC) derive from the Gondwanan continental crust, representing fragments of the upper plate during the Neo-Tethys subduction.
2. Three metamorphic stages and a prograde clockwise P–T trajectory are reconstructed for the NSMC eclogites: (a) an initial pre-eclogite stage preserved in garnet porphyroblasts, (b) a HP metamorphic stage at 1.90–2.15 GPa and 550–600 °C in the presence of Na-amphibole + zoisite ± carbonates and (c) a subsequent epidote-amphibolite-facies retrogression.
3.  $^{40}\text{Ar}/^{39}\text{Ar}$  dating of the eclogites constrains the minimum age of peak pressure conditions at  $191.14 \pm 0.50$  Ma, with the amphibolite-facies retrogression that was achieved at  $143.96 \pm 5.82$  Ma. These new data indicate that the onset of Neo-Tethys subduction began at least 7 Myrs earlier than what was previously reported.
4. Due to their upper plate derivation and age of metamorphism, we propose that the NSMC eclogites of the SSZ formed in response to tectonic erosion of the Jurassic active margin of Iran, with the Neo-Tethys subduction that started just after the closure of the Palaeo-Tethys suture zone to the north in response of a wide plate reorganization in the Iranian area following the Eo-Cimmerian orogeny.

**Supplementary material.** The supplementary material for this article can be found at <https://doi.org/10.1017/S0016756824000098>

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