Geochemistry and new zircon U–Pb geochronology of Mesoproterozoic Punugodu granite pluton, SE India: implications for anorogenic magmatism along the western margin of Nellore Schist Belt, India

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Abstract

We report a new zircon U–Pb age of 1257 ± 6 Ma for the Punugodu granite (PG) pluton in the Eastern Dharwar Craton (EDC), Southern India. The Mesoproterozoic PG is an alkali feldspar hypersolvus granite emplaced at shallow crustal level, as evident from the presence of rhyodacite xenoliths and hornfelsic texture developed in the metavolcanic country rocks of the Neoarchaean Nellore Schist Belt (NSB). Geochemically, the PG is metaluminous, ferroan and alkali-calcic, and is characterized by high SiO2 and Na2O + K2O, Ga/Al ratios >2.6, high-field-strength elements and rare earth element (REE) contents with low CaO, MgO and Sr, indicating its similarity to anorogenic, alkali (A-type) granite. The highly fractionated REE patterns with negative europium anomalies of PG reflect its evolved nature and feldspar fractionation. Mafic (MME) to hybrid (HME) microgranular enclaves represent distinct batches of mantle-derived magmas that interacted, mingled and undercooled within the partly crystalline PG host magma. Felsic microgranular enclaves (FME) having similar mineralogical and geochemical characteristics to the host PG most likely represent fragments of marginal rock facies of the PG pluton. The PG appears to be formed from an oceanic island basalt (OIB)-like source in an anorogenic, within-plate setting. The emplacement of PG (c. 1257 Ma) in the vicinity of Mesoproterozoic Kanigiri Ophiolite (c. 1334 Ma) shows an age gap of nearly 77 Ma, which probably suggests PG emplacement in an extensional environment along a terrain boundary at the western margin of the Neoarchaean NSB in the EDC.

1. Introduction

Globally, A-type granites occur in Precambrian and Phanerozoic terrains, mainly in extensional tectonic settings (e.g., Loiselle & Wones, 1979; Eby, 1990, 1992; Black & Liégeois, 1993). A-type granites are distinct from other granite types and are defined as relatively anhydrous with high SiO2 and Na2O + K2O, low MgO and high incompatible trace element contents including REE, Zr, Y, Nb and Ta (e.g., Eby, 1992; Frost et al. 1999; Bonin, 2007; Dall’Agnol et al. 2012; Li et al. 2014). A-type granites predominantly range from single-feldspar (hypersolvus) to two-feldspar (subsolvus) granites. In the hypersolvus varieties, the granitic rocks are composed of quartz and alkali feldspar; the latter yields coarse perthitic exsolution (e.g. Turner et al. 1992).

Several mechanisms have been proposed to explain the origin of A-type magmas: (1) direct fractionation of mantle-derived alkaline basals (e.g., Turner et al. 1992; Litvinovsky et al. 2002; Mushkin et al. 2003); (2) low degrees of partial melting of F- and Cl-enriched dry, lower-crustal granulitic residue from which a granite melt was previously extracted (e.g., Collins et al. 1982; Clemens et al. 1986; King et al. 1997); (3) low-pressure melting of calc-alkaline rocks at upper crustal levels (Skjerlie & Johnston, 1993; Patiño Douce, 1997); and (4) hybridization between anatexic granite and mantle-derived mafic magmas (Bedard, 1990; Kerr & Fryer, 1993; Wickham et al. 1996; Mingram et al. 2000).

Although fine-grained, dark-coloured microgranular enclaves are most common in calc-alkaline, metaluminous (I-type) granites (e.g. Didier & Barbarin, 1991; Kumar and Rino, 2006), they also occur in A-type granites and provide vital clues on the processes that have acted at different levels during the magma evolutionary history (Bonin, 1991). Further, microgranular enclaves and their distribution in felsic plutons, along with their textural and contact
relationships with felsic host, may provide insights into the magma chamber processes and dynamics (e.g. Vernon, 1984; Vernon et al., 1988; Kumar, 1995, 2010, 2020; Perugini et al., 2007).

In Peninsular India, A-type granites are widely distributed along the marginal zone of the Eastern Ghats Granulite Belt (EGGB), in the Southern Granulite Terrain, Mahakoshal Supracrustal Belt, Central India; the Mayurbhanj area, north of the Bastar Craton; and the Malani Igneous Province (e.g. Eby & Kochhar, 1990; Meert et al., 2010; Sarvothaman & Sesha Sai, 2010; Kumar et al., 2020; Yadav et al., 2020). In Southern India, the western margin of the Neoarchaean Nellore Schist Belt (NSB) is characterized by the emplacement of several granite plutons of Proterozoic age (Gupta et al., 1984; Dobmeier et al., 2006; Sesha Sai, 2013). These granite plutons form part of the Proterozoic granite emplacement zone located close to a major terrain boundary in the Eastern Dharwar Craton (EDC), i.e. the eastern margin of the Proterozoic Cuddapah Basin and Neoarchaean NSB. The Punugodu granite (PG) is one among these granite plutons spatially associated with other granites, gabbros, felsic volcanics and ophiolite mélangé rocks. The origin and evolution of PG have not yet been described. The present study therefore aimed to carry out a detailed investigation involving the field relation, petrography, geochemistry and geochronology of the PG occurring to the east of Cuddapah Basin, in order to characterize and discuss its petrogenesis, which may contribute to our understanding of the Proterozoic accretionary and rifting history of this region.

2. Geological setting

The Neoarchaean NSB constituted by metavolcanics and metasedimentary rocks is characterized by emplacement of mafic and felsic magmatic rocks of Proterozoic age (e.g. Vasudevan & Rao, 1976; Gupta et al., 1984; Sriniwasan & Roop Kumar, 1995; Raman & Murty, 1997; Dharma Rao & Reddy, 2007; Ravikant, 2010; Sesha Sai et al., 2013; Saha et al., 2015). The two domains that constitute the NSB are the upper western Udaigiri and the lower eastern Vinjamuru domains (Dobmeier & Raith, 2003; Saha et al., 2015). The mafic and felsic alkaline plutons of the NSB fall within the southern continuity of the Prakasam Alkaline Province (PAPF) (Ratnakar & Leelanandam, 1985; Rao et al., 1988; Ratnakar & Vijaya Kumar, 1985; Leelanandam et al., 2006). Several collision- and extension-related Proterozoic magmatic events played a significant role in the crustal accretion and growth to the east of Cuddapah Basin in the EDC (Divakara Rao et al., 1999; French et al., 2008; Dharma Rao et al., 2011; Sain et al., 2017; Sesha Sai, 2019).

The occurrence of ophiolite affinity rocks at Kandra (Kandra Ophiolite Complex) in the southern part of the NSB indicates that subduction would have initiated ~1.85 Ga (Sesa Sai, 2009; Vijaya Kumar et al., 2010). The eastern margin of the EDC evolved as the site for collision with the Napier Complex forming the 1.7 Ga old orogen (Chaudhuri et al., 2002; Dobmeier & Raith, 2003; Ravikant, 2010; Vijaya Kumar et al., 2010), resulting in the formation of granulites (1700 Ma), anorthosite (1690–1630 Ma) and granites (1650–1450 Ma) in the Ongole domain (Kovach et al., 2001; Simmat & Raith, 2008; Dharma Rao et al., 2012). The overlapping ages of these rocks indicates that the domain was magmatically active between at least c. 1720 and 1570 Ma (Henderson et al., 2014).

The plutons emplaced in the northern part of the NSB in the PAKP yield the zircon U–Pb Mesoproterozoic ages of Elchuru nepheline syenite (1321 ± 17 Ma; Upadhyay et al., 2006), Purimetla gabbro (1334 ± 15 Ma; Subramanyam et al., 2016), Uppalapadu nepheline syenite (1356 ± 41 Ma; Vijaya Kumar et al., 2007), Errakonda ferro syenite (1352 ± 2 Ma; Vijaya Kumar et al., 2007) and Kanigiri gabbro (1338 ± 27 Ma; Subramanyam et al., 2016), which are emplaced between the Udaigiri and Vinjamuru domains. The Kanigiri granite (1284 ± 4 Ma; Sain et al., 2017), Punugodu granite (PG) (1256 ± 6 Ma; present study) and Peddacharlapalle gabbro (1251.2 ± 9.4 Ma; Subramanyam et al., 2016) are ~100 Ma younger than the PAK rocks. The 1334 ± 20 Ma age of Kanigiri Ophiolite Mélangé and the age of the plutons emplaced in its vicinity show a gap of ~80 Ma. However, further studies are required on undated southern and northern plutons of the NSB and PAKP to shed light on the tectono-magmatic evolution of the NSB.

3. Field relationships and petrography

The PG pluton, covering an area of 7 km², occurs as a N–S-trending oval-shaped body commonly exposed to the east of Kanigiri granite and south of the Podili granite plutons (Fig. 1). Xenoliths of the pre-existing NSB volcanics are found hosted in the PG. These xenoliths are felsic, angular and fractured, and show sharp to partly reactive contacts with the host PG (Fig. 2a). In places, calc-silicate bands (Fig. 2b) of the pre-existing NSB are also observed. Reddy & Sesha Sai (2003) reported a rhodacitic (metavolcanic) rock having intimate contact with the PG. Along the contact with PG the hornfelsic textures are developed (Fig. 2c), which clearly indicate intrusive contact formed by thermal metamorphism. Therefore, the rhodacite represents pre-existing supracrustal NSB rock, now engulfed into the PG, and formed the large xenolith. In the southern margin of PG, contact with the chloride schist of NSB is observed (Fig. 2d). Due to soil development and the weathered contact of PG and chloride schist of NSB, the actual nature of the contact is unknown.

The PG hosts abundant microgranular enclaves (ME), These ME are fine-grained, mafic–melanocratic (MME) to porphyritic (hybrid) mesocratic (HME), and vary in size from a few mm to metres, with rounded, elliptical, spherical and lenticular shapes commonly having sharp, crenulated and occasionally diffuse contacts with the host PG (Fig. 2e–h). These field features of ME indicate ME magma mingling and undercooling within the partly crystalline PG melt (e.g. Kumar et al., 2004). The HME exhibit characteristics that are intermediate between MME and felsic microgranular enclaves (FME) (Fig. 2f). The N–S-oriented nature of ME perhaps indicates the phenomenon of magmatic flowage attained during the emplacement and dynamics of coexisting ME and PG magmas (Fig. 2g) prior to solidification of the entire system as described by Kumar (2020). The FME are fine-grained, linear to spherical to lenticular (Fig. 2h), most likely representing fragments of early, chilled margin facies of PG pluton and disrupted as FME due to prevailing convective force in the PG chamber (e.g. Didier & Barbarin, 1991; Kumar, 2010). The depth of emplacement is one factor that plays a vital role in the crystallization of A-type alkalifeldspar granites. Felsic magmas of A-type affinity are products of residual liquids resulting from the differentiation phenomenon in the magma chambers with varying depths (Bonin, 1998).

A total of 29 rock samples, comprising 14 PG, four xenoliths, four FME, three HME and four MME samples, were collected during fieldwork covering the outcrops of PG pluton. Sample locations are shown in Figure 1.
3. Petrography of PG

Detailed petrographic studies reveal that the PG is essentially composed of relatively large subhedral laths of perthitic K-feldspar and quartz, while the mafic phases noticed are biotite and amphibole, with zircon, titanite, fluorite, apatite and opaques as the accessories. Relict clinopyroxene is also noticed in places in...
Fig. 2. (Colour online) Field photographs showing the field relationship of the PG pluton. (a) Xenolith with fractures showing sharp contact with host rock. (b) Occurrence of a calc-silicate xenolith of NSB in PG. (c) Presence of hornfelsic rock at the contact with pre-existing metavolcanic rock. (d) Field photograph showing contact between PG and chlorite schist of NSB country rock. (e) Fine-grained, mafic microgranular enclave surrounded by mafic clusters. (f) Hybrid microgranular enclave with ellipsoidal shape. (g) Field photograph showing oriented nature of magmatic enclaves at the pluton contact indicating magma mingling. (h) Fine-grained felsic microgranular enclave with rounded shape.
the PG. Amphibole is pleochroic in shades of brown to blue. The presence of single feldspar in the form of microcline perthite indicates the hypersolvus nature of PG (Fig. 3a and b). Petrographic studies of the PG samples collected along the margins of the pluton reveal that quartz is anhedral and exhibits wavy extinction and recrystallization indicating a deformational regime (Fig. 3c). Perlitic K-feldspars are characteristically twinned (Fig. 3d). Perlitic occurs as string, vein and patchy types (Anderson, 1928; Smithson, 1963). String perlitic are formed due to exsolution, as evident from small, well-oriented blebs perlitic, whereas the patchy and coarse vein types of perlitic are formed due to replacement evident from irregular, redistributed and enlarged albite blebs, which occur close to and on the margin of the grain (e.g. Alling, 1938). Fluorite is observed, associated with biotite (Fig. 3e). Bluish amphibole occurs as subhedral prismatic grains in places (Fig. 3f). Clusters of mafic phases represented mainly by amphiboles and associated opaques are observed in parts of the quartz-feldspathic material (Fig. 3g). Fine-grained skeletal opaques are observed in the vicinity of biotite and amphibole. Zircon is euhedral and exhibits zoning (Fig. 3h).

3.b. Petrography of enclaves

Petrographic studies indicated variation in the mineralogy of three types of magmatic enclaves observed in PG, namely FME, HME and MME. The FMEs are mineralogically similar to host granite and mainly consist of quartz, microcline, plagoclase and biotite (Fig. 4a and b). The HMEs are fine-grained and porphyritic, exhibiting typical igneous texture with felsic phenocrysts (xenocrysts). Quartz xenocrysts (ocelli) with biotite and hornblende-rich mantle (Fig. 4c), perlitic and opaques with biotite rim and clusters are common (Fig. 4d). Mineralogically, the MMEs consist of plagioclase and pyroxene as essential minerals and exhibit subhedral texture (Fig. 4e). Twinned plagioclase crystals occur as elongated laths in places in the MMEs, a textural feature suggesting magmatic origin (Fig. 4 f). Biotite appears as interstitial flakes in the MMEs. Subhedral grains of Fe-oxides occur as accessory minerals in the MMEs (Fig. 4g). Most MMEs show diffusive and cuspathe contacts with host PG, indicating their magmatic nature. This type of enclave contact mainly appears in some particular cases where large volumes of mafic magma dominate over felsic magma (Barbarin & Didier, 1991). The enclaves as xenoliths represent the caught-up patches of the NSB metavolcanics and hence are different from the observed ME types. Xenoliths show feeble deformation, which is evident from the recrystallized quartz and distorted twin lamellae in plagioclase. Petrographic studies show the presence in the metavolcanic xenolith of quartz porphyroclasts along with feldspar, hornblende and biotite (Fig. 4h).

4. Sampling and analytical techniques

Relatively fresh and unweathered samples were collected during the fieldwork. High-quality thin-sections were made for the systematic petrographic studies. Rock samples for major, trace and rare earth elemental analysis were crushed and pulverized to fine powders manually using the agate mortar and pestle.

4.a. Major oxides and trace elements

Geochemical analyses were carried out at the Geochemistry Division, CSIR – National Geophysical Research Institute (NGRI), Hyderabad, India. Major-element abundances were determined on pressed powder pellets using X-Ray Fluorescence (XRF) (Phillips MAGIX PRO Model 2440) following the methods outlined by Krishna et al. (2007). Trace and rare earth elements were determined by high-resolution inductively coupled mass spectrometer (HR-ICP-MS; Nu Instruments Attom, UK) in jump wiggle mode. The analytical procedure, precision and accuracy for HR-ICP-MS are reported in Manikyamba et al. (2016). ¹⁰³Rh was used as an internal standard, and external drift was corrected by repeated analyses of standards used for calibration.

4.b. Zircon separation and U–Pb analysis

Zircon grain extractions were carried out after crushing the rock samples into ~80–250 μm sieves fractions and were processed on the Wilfley shake table to separate heavy mineral fractions. The heavies were then subjected to isodynamic magnetic separation before handpicking zircons under a binocular microscope. Zircon grains were mounted in 25 mm epoxy resin blocks, diamond-polished, carbon-coated and cathodoluminescence (CL)-imaged on a TESCAN LMH3 EM mounted with a Rainbow colour CL detector. Zircon U–Pb dating was performed using Thermo X SeriesICP-MS coupled to a New Wave Universal Platform 213 nm Nd–YAG laser ablation system with a large-format ablation cell operated with a combination of He and Ar gases as detailed in Babu et al. (2009). The data were processed offline using ‘IsoplotR’ (Vermeesch, 2018) and ‘Glitter software’ to compute the ages and corrected for common Pb using Andersen’s scheme (Andersen, 2002).

Results

The major, trace and rare earth element data of the granite samples, along with those of the xenoliths, FME, HME and MME analysed in the present study, are presented in Table 1.

5.a. Geochemistry of PG and enclaves

Major-oxide geochemistry of PG reveals high SiO₂ (avg. 70.41 wt %) and high Na₂O + K₂O (avg. 9.40 wt %) contents, a feature that corresponds to the anorogenic, alkalai (A-type) granites (Whalen et al. 1987). The SiO₂ concentrations in enclaves are as follows: in xenoliths from 61.2 to 64.9 wt %, FME from 66.5 to 67.4 wt %, HME from 61.2 to 64.9 wt %, and MME from 66.5 to 67.4 wt %. The K₂O + Na₂O content in xenoliths, FME, HME and MME ranges from 8.58 to 9.48 wt %, 8.27 to 9.5 wt %, 8.34 to 9.25 wt % and 3.74 to 5.24 wt % respectively. A detailed account of major oxide, trace element and rare earth element geochemical characteristics of PG pluton was presented previously (Narshimha et al. 2018). In the MALI (Na₂O + K₂O–CaO) vs silica diagram (Frost et al., 2001), the PG granite, FME and xenoliths exhibit an alkaline to alkali-calcic trend (Fig. 5a). The MMEs in the FeO* vs silica diagram (Frost & Frost, 2008) occupy the tholeiitic and alkaline fields (Fig. 5b). The Fe number of 13 PG samples varies from 0.76 to 0.97, indicating its strong ferron character. In the A/NK – A/CNK alumina saturation diagram (Fig. 5c), the majority of PG samples fall in the metaluminous domain, while a few fall close to the boundary of the metaluminous and peralkaline fields (Shand, 1943). In the normative anorthite (An) – albite (Ab) – orthoclase (Or) diagram (O’Connor, 1965), the samples fall in the granite field (Fig. 5d). In the R₁–R₂ multicationic parameters (De la Roche et al. 1980) based tectonic classification scheme (after Batchelor & Bowden, 1985), the PG shows affinity with syn-collision to late orogenic tectonic settings (Fig. 5e). Pearce et al. (1984) classified the granites into a within-plate, collisional,
Fig. 3. (Colour online) (a) Photomicrograph under crossed Nicols showing coarse-grained perthitic feldspar. (b) Photomicrograph of microcline microperthite with biotite cluster. (c) Admixture of recrystallized quartz along with the feldspar grain boundary indicating recrystallization due to deformation. (d) Photomicrograph under crossed Nicols showing Carlsbad twinning in perthitic K-feldspar. (e) Photomicrograph in plane-polarized light showing fluorite associated with biotite. (f) Photomicrograph in plane-polarized light showing relict clinopyroxene surrounded by alkali amphibole. (g) Photomicrograph in plane-polarized light showing skeletal opaques in the vicinity of mafic minerals. (h) Zoned euhedral zircon (plane polarized light). Mineral abbreviations used as in Whitney & Evans [2010].
Fig. 4. (Colour online) (a, b) Photomicrographs under crossed Nicols showing quartz, plagioclase, microcline and biotite minerals in FME. (c) Photomicrograph under plain polarized light showing quartz xenocrysts with biotite- and hornblende-rich mantle in HME. (d) Photomicrographs under crossed Nicols showing perthites and clusters of biotite in HME. (e, f) Photomicrograph showing plagioclase and pyroxene with sub-ophitic texture in MME. (g) Photomicrograph showing the cluster of opaques in MME. (h) Photomicrograph under crossed Nicols showing quartz porphyroclast in NSB metavolcanic xenolith.
| Sl. No. | 1  | 2  | 3  | 4  | 5  | 6  | 7  | 8  | 9  | 10 | 11 | 12 | 13 | 14 |
|--------|----|----|----|----|----|----|----|----|----|----|----|----|----|----|
|        |     | PG-1 | PG-2 | PG-3 | PG-4 | PG-5 | PG-6 | PG-7 | PG-8 | PG-9 | PG-10 | PG-11 | PG-12 | PG-13 | PG-14 |
| **Granite** |    |    |     |     |     |     |     |     |     |     |     |     |     |     |
|         |    |     |     |     |     |     |     |     |     |     |     |     |     |     |     |
| **Major elements (wt %)** |    |     |     |     |     |     |     |     |     |     |     |     |     |     |     |
| SiO₂   | 71.2 | 64.9 | 71.2 | 69.8 | 71.4 | 69.6 | 72.7 | 69.5 | 68.9 | 70.2 | 73.1 | 71.8 | 71.3 | 70.2 |
| Al₂O₃  | 13.3 | 13.9 | 12.5 | 13.2 | 13.4 | 13.3 | 11.7 | 14.5 | 13.8 | 13.3 | 12.0 | 13.7 | 13.7 | 14.2 |
| Fe₂O₃  | 3.6  | 4.6  | 2.9  | 3.6  | 3.2  | 3.4  | 3.0  | 3.6  | 3.0  | 3.2  | 2.3  | 2.7  | 2.9  |       |
| MnO    | 0.1  | 0.1  | 0.1  | 0.1  | 0.1  | 0.1  | 0.1  | 0.1  | 0.1  | 0.1  | 0.0  | 0.1  | 0.1  |       |
| MgO    | 0.1  | 3.6  | 1.1  | 0.5  | 0.3  | 0.8  | 0.6  | 0.2  | 0.9  | 0.6  | 0.3  | 0.3  | 0.2  | 0.9  |
| CaO    | 1.0  | 3.1  | 1.2  | 1.2  | 1.0  | 0.9  | 1.3  | 1.1  | 0.7  | 1.0  | 0.8  | 1.1  |       |       |
| Na₂O   | 3.3  | 4.0  | 3.5  | 3.7  | 3.4  | 3.7  | 3.6  | 3.8  | 3.5  | 3.3  | 3.6  | 3.4  | 3.8  | 3.6  |
| K₂O    | 6.0  | 3.8  | 5.9  | 6.3  | 5.8  | 6.4  | 5.7  | 6.4  | 5.6  | 6.4  | 5.7  | 6.2  | 5.7  |       |
| TiO₂   | 0.4  | 0.5  | 0.3  | 0.4  | 0.3  | 0.3  | 0.3  | 0.3  | 0.3  | 0.3  | 0.3  | 0.2  | 0.3  | 0.3  |
| P₂O₅   | 0.0  | 0.1  | 0.1  | 0.0  | 0.0  | 0.0  | 0.0  | 0.1  | 0.0  | 0.0  | 0.0  | 0.0  | 0.0  | 0.0  |
| LOI    | 0.6  | 1.1  | 0.7  | 0.8  | 0.5  | 0.7  | 0.5  | 1.2  | 0.8  | 1.1  | 0.6  | 0.8  | 0.8  | 0.3  |
| SUM    | 99.6 | 99.7 | 99.4 | 99.4 | 99.2 | 99.1 | 99.8 | 98.8 | 99.4 | 99.4 | 99.2 | 99.7 | 99.1 |       |
| Fe⁺    | 0.97 | 0.56 | 0.72 | 0.87 | 0.91 | 0.81 | 0.83 | 0.93 | 0.80 | 0.83 | 0.88 | 0.88 | 0.99 | 0.76 |
| **Trace elements (ppm)** |    |     |     |     |     |     |     |     |     |     |     |     |     |     |
| La     | 312.8 | 101.4 | 242.5 | 320.0 | 187.9 | 161.3 | 144.1 | 160.5 | 191.3 | 179.2 | 170.0 | 215.0 | 225.7 | 170.9 |
| Ce     | 1056.7 | 190.2 | 469.8 | 1047.8 | 360.0 | 314.8 | 280.4 | 314.3 | 373.7 | 336.1 | 326.9 | 427.4 | 439.4 | 332.3 |
| Pr     | 64.9 | 19.2 | 49.9 | 62.2 | 36.8 | 32.9 | 29.2 | 33.2 | 39.3 | 34.6 | 35.1 | 43.5 | 46.0 | 34.5 |
| Nd     | 238.7 | 69.4 | 184.6 | 224.2 | 133.3 | 119.4 | 105.8 | 120.9 | 144.0 | 125.5 | 129.7 | 149.9 | 167.2 | 124.8 |
| Sm     | 46.3 | 13.1 | 35.5 | 39.4 | 25.7 | 24.0 | 21.6 | 24.0 | 28.5 | 23.7 | 26.8 | 27.7 | 32.5 | 24.7 |
| Eu     | 0.5 | 0.8 | 0.5 | 0.6 | 0.5 | 0.5 | 0.4 | 0.5 | 0.6 | 0.5 | 0.6 | 0.5 | 0.4 | 0.5 |
| Gd     | 38.3 | 11.4 | 29.2 | 32.1 | 22.1 | 20.7 | 18.8 | 20.4 | 24.5 | 21.2 | 23.5 | 24.7 | 27.6 | 21.7 |
| Tb     | 5.7 | 1.9 | 4.3 | 4.6 | 3.6 | 3.5 | 3.2 | 3.4 | 4.0 | 3.4 | 4.0 | 4.4 | 4.4 | 3.7 |
| Dy     | 34.8 | 13.5 | 25.2 | 27.4 | 24.4 | 23.9 | 21.7 | 22.5 | 26.2 | 22.4 | 27.7 | 32.0 | 29.1 | 24.8 |
| Ho     | 3.6 | 1.6 | 2.6 | 2.8 | 2.8 | 2.8 | 2.5 | 2.5 | 2.9 | 2.6 | 3.2 | 3.9 | 3.2 | 2.8 |
| Er     | 11.2 | 5.5 | 7.7 | 9.2 | 9.0 | 8.2 | 8.4 | 9.4 | 8.3 | 10.2 | 13.2 | 10.6 | 9.2 |
| Tm     | 1.3 | 0.7 | 0.9 | 1.1 | 1.1 | 1.2 | 1.0 | 1.0 | 1.1 | 1.0 | 1.3 | 1.7 | 1.3 | 1.2 |
| Yb     | 12.6 | 7.7 | 8.5 | 11.1 | 10.8 | 11.3 | 10.0 | 10.1 | 11.1 | 10.1 | 11.8 | 15.8 | 12.8 | 11.2 |
| Lu     | 2.0 | 1.3 | 1.4 | 1.8 | 1.7 | 1.8 | 1.5 | 1.6 | 1.8 | 1.6 | 1.9 | 2.3 | 2.0 | 1.7 |
| Sc     | 1.8 | 4.7 | 1.8 | 1.9 | 1.9 | 1.9 | 2.0 | 2.1 | 2.3 | 2.0 | 2.2 | 2.4 | 2.1 |       |

(Continued)
| Sl. No. | 1  | 2  | 3  | 4  | 5  | 6  | 7  | 8  | 9  | 10 | 11 | 12 | 13 | 14 |
|--------|----|----|----|----|----|----|----|----|----|----|----|----|----|----|
| **Granite** |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Sample | PG-1 | PG-2 | PG-3 | PG-4 | PG-5 | PG-6 | PG-7 | PG-8 | PG-9 | PG-10 | PG-11 | PG-12 | PG-13 | PG-14 |
| V      | 2.0 | 7.9 | 1.4 | 1.6 | 1.3 | 1.3 | 1.2 | 1.2 | 1.1 | 1.6 | 1.1 | 1.4 | 1.2 | 1.4 |
| Cr     | 7.9 | 8.2 | 7.9 | 8.1 | 8.2 | 8.1 | 8.3 | 8.1 | 8.1 | 8.1 | 8.0 | 8.3 | 8.0 | 8.3 |
| Co     | 0.4 | 8.8 | 0.4 | 0.5 | 0.5 | 0.5 | 0.6 | 0.5 | 0.6 | 1.2 | 0.8 | 1.0 | 0.5 | 0.8 |
| Ni     | 1.5 | 4.1 | 1.4 | 1.3 | 1.2 | 1.2 | 1.4 | 1.2 | 1.4 | 1.5 | 1.7 | 1.3 | 1.2 | 1.5 |
| Cu     | 0.5 | 0.7 | 0.5 | 0.5 | 0.5 | 0.5 | 0.4 | 0.5 | 0.5 | 0.5 | 0.4 | 0.5 | 0.5 | 0.5 |
| Zn     | 42.3 | 27.6 | 37.6 | 48.9 | 45.1 | 52.4 | 38.8 | 44.7 | 44.5 | 38.4 | 44.0 | 21.0 | 30.2 | 45.8 |
| Ga     | 37.6 | 26.9 | 37.0 | 38.9 | 36.9 | 40.4 | 35.4 | 40.1 | 38.3 | 33.9 | 36.0 | 36.0 | 37.6 | 38.1 |
| Rb     | 197.3 | 136.4 | 179.1 | 224.7 | 241.2 | 251.5 | 225.3 | 247.3 | 241.0 | 212.8 | 230.6 | 345.2 | 278.6 | 223.2 |
| Sr     | 9.6 | 93.6 | 10.8 | 10.4 | 11.4 | 9.9 | 8.4 | 13.0 | 12.6 | 17.8 | 12.3 | 15.0 | 10.7 | 14.3 |
| Y      | 156.6 | 79.6 | 108.1 | 129.5 | 137.9 | 137.5 | 123.6 | 125.6 | 141.8 | 129.0 | 151.0 | 201.8 | 144.7 | 141.5 |
| Zr     | 128.4 | 341.8 | 85.5 | 153.6 | 279.0 | 311.8 | 275.6 | 274.4 | 282.0 | 255.8 | 296.2 | 265.4 | 113.7 | 356.1 |
| Nb     | 216.6 | 99.3 | 131.6 | 194.9 | 183.0 | 188.1 | 167.1 | 163.0 | 173.1 | 184.9 | 165.7 | 211.4 | 190.1 | 180.0 |
| Cs     | 1.6 | 1.9 | 1.5 | 2.0 | 1.9 | 2.0 | 1.6 | 1.8 | 2.4 | 1.6 | 2.7 | 2.6 | 2.6 | 3.1 |
| Ba     | 32.7 | 243.5 | 40.0 | 37.4 | 58.4 | 56.4 | 69.5 | 69.9 | 72.5 | 132.3 | 131.9 | 96.9 | 45.2 | 89.3 |
| Hf     | 4.9 | 11.8 | 3.2 | 6.1 | 10.2 | 12.9 | 10.1 | 10.2 | 10.3 | 8.2 | 11.3 | 11.9 | 4.6 | 12.7 |
| Ta     | 8.2 | 7.4 | 4.7 | 8.2 | 8.1 | 11.3 | 8.5 | 8.1 | 8.0 | 8.5 | 8.1 | 12.1 | 9.0 | 9.0 |
| Pb     | 17.4 | 23.6 | 16.9 | 17.3 | 21.5 | 19.8 | 14.5 | 19.6 | 14.8 | 35.7 | 17.9 | 16.0 | 18.4 | 16.6 |
| Th     | 51.6 | 35.1 | 36.4 | 47.1 | 43.0 | 45.0 | 39.2 | 38.7 | 40.4 | 41.0 | 41.9 | 118.4 | 55.5 | 44.5 |
| U      | 6.0 | 10.4 | 3.8 | 5.2 | 6.4 | 6.4 | 8.4 | 6.2 | 5.9 | 5.0 | 4.2 | 20.6 | 8.0 | 8.0 |

| Sl. no. | 15 | 16 | 17 | 18 | 19 | 20 | 21 | 22 | 23 | 24 | 25 | 26 | 27 | 28 | 29 |
|---------|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|
| **Enclave** |    |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| Sample | Xenolith-1 | Xenolith-2 | Xenolith-3 | Xenolith-4 | HME-1 | HME-2 | HME-3 | FME-1 | FME-2 | FME-3 | FME-4 | MME-1 | MME-2 | MME-3 | MME-4 |
| Major elements (wt %) |    |    |    |    |    |    |    |    |    |    |    |    |    |    |    |
| SiO₂ | 64.9 | 64.2 | 61.5 | 61.2 | 53.3 | 49.4 | 62.9 | 66.5 | 67.4 | 66.9 | 66.9 | 45.2 | 48.7 | 49.0 | 47.6 |
| Al₂O₃ | 13.2 | 13.5 | 13.2 | 13.1 | 14.5 | 11.0 | 13.8 | 13.7 | 13.6 | 14.0 | 13.9 | 13.9 | 13.1 | 13.2 | 13.4 |
| Fe₂O₃ | 6.4 | 6.1 | 7.3 | 7.9 | 8.2 | 10.7 | 6.7 | 6.8 | 6.1 | 4.3 | 5.3 | 11.0 | 12.0 | 11.5 | 11.5 |
| MnO | 0.1 | 0.1 | 0.1 | 0.1 | 0.1 | 0.2 | 0.1 | 0.1 | 0.1 | 0.1 | 0.1 | 0.1 | 0.1 | 0.1 | 0.1 |
| MgO | 0.8 | 1.3 | 2.9 | 3.5 | 2.9 | 9.9 | 0.9 | 0.4 | 0.2 | 0.5 | 0.5 | 12.3 | 7.6 | 8.2 | 9.4 |
| CaO | 3.1 | 3.5 | 3.6 | 3.5 | 1.6 | 6.7 | 4.0 | 2.0 | 1.7 | 2.7 | 2.0 | 9.6 | 9.7 | 10.8 | 10.0 |
| Element   | Na2O | K2O | TiO2 | P2O5 | SUM | Trace elements (ppm) |
|-----------|------|-----|------|------|-----|---------------------|
|           |      |     |      |      |     | La                  |
|           |      |     |      |      |     | Ce                  |
|           |      |     |      |      |     | Pr                  |
|           |      |     |      |      |     | Nd                  |
|           |      |     |      |      |     | Sm                  |
|           |      |     |      |      |     | Eu                  |
|           |      |     |      |      |     | Gd                  |
|           |      |     |      |      |     | Tb                  |
|           |      |     |      |      |     | Dy                  |
|           |      |     |      |      |     | Ho                  |
|           |      |     |      |      |     | Er                  |
|           |      |     |      |      |     | Tm                  |
|           |      |     |      |      |     | Yb                  |
|           |      |     |      |      |     | Lu                  |
|           |      |     |      |      |     | Sc                  |
|           |      |     |      |      |     | V                   |
|           |      |     |      |      |     | Cr                  |
|           |      |     |      |      |     | Co                  |
|           |      |     |      |      |     | Ni                  |
|           |      |     |      |      |     | Cu                  |
|           |      |     |      |      |     | Zn                  |
|           |      |     |      |      |     | Ga                  |
|           |      |     |      |      |     | Rb                  |
|           |      |     |      |      |     | Sr                  |
|           |      |     |      |      |     | Y                   |
|           |      |     |      |      |     | Zr                  |

(Continued)
volcanic arc and oceanic-ridge granites mostly based on the content of high-field-strength elements (HFSE). The within-plate origin of the PG samples is substantiated by the Rb vs Y + Nb plot (Fig. 5f). The Rb vs Y + Nb plot (Fig. 5f) of PG and enclaves shows that MMEs are alkaline occupying the basaltic field (Fig. 5g). The HMEs show mafic-felsic intermediate compositions occupying the trachyte-trachyandesite and phonotephrite fields.

The Harker variation diagrams (Fig. 6) show the variation of major element oxides with silica as the magma differentiate in which Na₂O and K₂O increase, indicating a positive correlation during the evolution of PG pluton. The MgO, FeO, TiO₂ and CaO decrease with increasing silica. Al₂O₃ remains almost constant or slightly increases with differentiation. The K₂O content in FME ranges from 4.3 to 3.6 wt %, HME from 4.9 to 3.2 wt %, and MME from 2.2 to 1.4 wt %, indicating an overall decrease in K₂O content from FMEs to MMEs. The high Fe₂O₃ content in MME ranges from 10.95 to 11.48 wt %, which can be ascribed to the presence of mafic minerals. The Rb concentrations in the PG samples are generally high and vary from 136 to 278 ppm. This is concomitant with the high abundance of K₂O in the granite. Ba, which can also substitute to K, is less abundant, ranging from 32 to 132 ppm. The decrease in Ba content with increasing SiO₂ (Fig. 6) indicates the fractionation of K-feldspar during the crystallization of PG. The Rb contents are moderate, varying between 136 and 278 ppm. The Ca and Sr contents show decreasing trends with increasing SiO₂. The granite samples are also characterized by moderately high concentrations of U (4.2 to 20.5 ppm) and Th (35 to 55.5 ppm) along with HFSE such as Zr (85.5 to 356 ppm), Nb (99 to 216 ppm) and Y (79.6 to 201.8 ppm), and low concentrations of transition metal trace elements Ni (1.1 to 4 ppm), Cr (7.8 to 8.2 ppm) and Co (0.3 to 1.1 ppm). Trace element geochemistry indicates that the MME consist of relatively high Ni (10.6 to 88 ppm), Cr (63 to 283 ppm) and Co (45 to 115 ppm). The HFSE contents of Zr (144 to 466 ppm) and Y (28 to 79 ppm) in MME show moderate abundances.

The PG samples are characterized by low to moderately fractionated REE patterns (Fig. 7a) with (La/Yb)N varying between 9.05 and 19.26. The PG samples are characterized by strong LREE fractionation ((La/Sm)N = 3.91–5.0) with flat HREE ((Gd/Yb)N = 0.7–3.3) patterns. The PG samples also exhibit negative Eu anomalies (Eu/Eu)ⁿ ranging from 0.04 to 0.07. The chondrite-normalized (after Nakamura, 1974) REE patterns of xenolith, FME and HME are enriched in LREE and show prominent negative Eu anomalies (Fig. 7b, c and d), indicating a significant role of plagioclase fractionation from the parent magma. The chondrite-normalized REE patterns of the MME exhibit marginally enriched LREE and relatively depleted HREE (Fig. 7e). Low-amplitude negative to no Eu anomaly is observed in the MME, with the (Eu/Eu)ⁿ ranging from 0.3 to 1.0. The primitive-mantle-normalized (after Sun & McDonough, 1989) multi-element diagram (Fig. 7f) of the PG samples shows prominent negative anomalies in Ba, Sr, P, Eu and Ti, while Th, U, Ce, Pr, Nb, Sm, Dy, Y and Nd show positive anomalies. The negative anomalies of Sr and Eu are related to the fractionation of plagioclase in the source region (Hanson, 1978). The negative anomalies of P and Ti reflect the fractionation ofapatite and Fe–Ti oxides. The overall geochemical characteristics of the granites from PG pluton indicate its anorogenic, alkaline (A-type) nature (Whalen et al., 1987). The incompatible trace-element-normalized (after Sun & McDonough, 1989) multi-element diagrams (Fig. 7g, h and i) for the xenolith, FME and HME depict negative Ba, Sr and Ti anomalies and...
Fig. 5. (Colour online) (a) MALI vs silica diagram (Frost et al., 2001) showing alkali-calcic nature of Punugodu granite. (b) FeO* vs silica diagram (Frost & Frost, 2008) showing MME occupy the tholeiitic/alkaline field. (c) A/NK vs A/CNK diagram (Shand, 1943; Maniar & Piccoli, 1989) for granites and enclaves. (d) Normative anorthite (An)–albite (Ab)–orthoclase (Or) diagram (O’Connor, 1965). (e) R1–R2 tectonic classification diagram (after Batchelor & Bowden, 1985) showing the position of Punugodu granite in syn-collision to late orogenic tectonic setting. R1 = 4Si – 11(Na + K) – 2(Fe + Ti); R2 = 6Ca + 2Mg + Al. (f) Rb vs (Y + Nb) diagram (after Pearce et al., 1984, 1996) showing the position of Punugodu granite in WPG. (g) Total-alkali silica (TAS) diagram (Le Maitre, 2002) of Punugodu samples. Note: The symbols denoting granite, xenoliths, FME, HME and FME are applicable for all geochemical plots.
positive Rb anomalies, whereas the MMEs show negative Ba with constant Sr and Ti and positive Rb anomaly (Fig. 7j).

5.b. Zircon U-Pb geochronology

The PG-8 granite sample with abundant zircons is chosen for U-Pb geochronology. Zircon grains extracted from this sample are well-developed euhedral grains, typically 80–200 μm long with aspect ratios of 2:1 to 3:1. They show concentric growth zoning typical of magmatic zircons (Corfu et al. 2003), and, in a few cases, convolute zoning suggests crystallization from melts. The CL images of representative zircon grains are given in Figure 8. The zircon grains show well-preserved prismatic terminations suggesting limited magmatic resorption. Most of the zircons have systematic growth zoning from core to rim. Luminescent outer rims are rare, suggesting a general lack of metamorphic overgrowth. Rarely, an effect of alteration is evident in the development of bright luminescent patches across growth zoning.

In general, the laser spots were placed in homogeneous interiors of zircons, avoiding older (inherited) domains and visibly altered segments. In situ laser-ablation ICP-MS U–Th–Pb isotopic data for individual laser spots in zircons from the sample are given in Table 2. Age calculations are discussed based on a $^{206}\text{Pb} / ^{238}\text{U}$ concordia diagram (Fig. 9). The concordant zircon grains have U content between 7.6 and 384 ppm with an average of 95.4 ppm, Th ranging from 5.6 ppm to 427.7 ppm with an average 75.6 ppm, and Th/U = 0.43 to 1.49, attesting to their typical magmatic character (Rubatto, 2002). The zircons yield a concordia upper intercept U–Pb age of 1257 ± 6 Ma (2σ errors), MSWD = 7.5. We interpret 1257 ± 6 Ma concordia upper intercept age as the time of emplacement and magmatic crystallization of the Punugodu granite pluton.

6. Discussion

6.a. Genetic type and tectonic setting

The Punugodu granite is mineralogically an alkali feldspar granite and is characterized by high SiO$_2$ and Na$_2$O + K$_2$O. The average Fe number for PG is 0.83, indicating its ferron nature. In the A/NK vs A/CNK diagram (Fig. 5c), the sample of PG granite falls at the juncture of metaluminous and peralkaline fields. Based on the major oxide geochemistry, alkali feldspar granites of ferroan affinity, metaluminous and peralkaline by virtue of their alumina saturation, have been recognized as alkalic (A-type) granites (e.g. Frost & Frost, 2011). Whalen et al. (1987) proposed several
discrimination diagrams to distinguish the A-type granite from other granite types such as I-, S- and M-types. The 10 000 $^*$ Ga/Al values vary with Zr, Nb and FeO/MgO content, and Na$_2$O $+$ K$_2$O can discriminate A-type granites. The separating line in the discrimination diagram is at Ga/Al $>$ 2.6. All the samples of the PG are positioned exclusively in the field of A-type granite in the Zr and Nb vs Ga/Al diagrams (Fig. 10a and b). The high concentration of Na$_2$O $+$ K$_2$O, high Fe number and enriched
HFSE (Zr, Nb and Y) contents along with enriched LREE in the PG correspond with A-type granites (Whalen et al., 1987; Eby, 1992). Eby (1992) classified the A-type granites into subtypes A1 and A2, based on the Y/Nb ratio. The A1 group (Y/Nb < 1.2) was interpreted as alkaline magma derived from an oceanic island basalt (OIB)-like source in a proper anorogenic within-plate setting. The PG shows Y/Nb ratios of <1.2, ranging from 0.6 to 0.91, indicating its A1-type granite, commonly emplaced in an extensional regime in an anorogenic within-plate setting. This is further corroborated by ternary Y–Nb–3Ga and Y–Nb–Zr/4 plots (Fig. 10c and d) for the PG samples (Eby, 1992).

Pearce et al. (1984), based on the compilation of >600 analyses of felsic rocks, divided the granites into four major tectonic groups: ocean ridge granite (ORG), volcanic arc granite (VAG), within-plate granite (WPG) and collisional granite (COLG). In the Rb vs Y + Nb graph, the PG clearly plots within the WPG field (Fig. 5f). This setting for some A-type granites has also been described from occurrences elsewhere in the world (e.g. Eby, 1992). The prominent positive anomalies in granite and enclave samples observed for Rb, Th, U, Ce, Pr, Nb, Sm, Dy, Y and Nd are typical of anorogenic-related granite magmatism.

6.b. Magma mingling and mixing

The presence of oval-, sub-oval- and elliptical-shaped ME hosted in the PG indicates the hybridizing magma system (Kumar et al. 2004). The host PG and the enclaves such as xenoliths and FME show a continuous variation in SiO₂ concentrations, producing tightly linear to slightly curved trends on Harker variation diagrams (Fig. 6), strong evidence for a genetic link between them. However, the observed wide composition of HME may be due to the chemical mixing between two contrasting magma end-members in varying proportions (Kumar & Rino, 2006). The fine-grained, igneous-textured mafic to hybrid ME from I- and A-type granites has been considered to represent mafic and/or hybrid magma components added to intermediate or felsic magma chambers (e.g. Holden et al. 1987; Vernon et al. 1988; Bedard, 1990; Didier & Barbarin, 1991; Bonin, 2004; Yang et al. 2004;
Kumar, 2020). Silica contents in the MEs indicate a broad bimodal distribution. The SiO2 contents in MME and in some HME range from 45 to 53 %, while the SiO2 contents in FME and granite samples show a relatively wide variation from 61 to 73 %, separated by a silica gap of c. 8 wt %. The relative enrichment of Ni and Cr in the MME and HME samples suggests that they might have formed from mantle-derived mafic and mafic–felsic (hybrid) magmas respectively, injected into the crystallizing host magma chamber as described elsewhere (Kumar et al., 2017). Notable variation in the Ba content in some of the MME samples is perhaps due to geochemical heterogeneities in thefeldspars during magma mixing (e.g. Bonin, 2004; Słaby et al., 2011).

6.c. Temperature and pressure

Zirconium saturation level (ppm) as a function of cationic ratio M = (Na + K + 2Ca)/(Al. Si) (Watson & Harrison 1983) shows two different ranges of crystallization temperatures for granite, 750–800°C and 850–900°C (Fig. 10e). This may be because of crystallization of zircon right from liquidus to solidus. Based on

| Analysis_# | Th ppm | U ppm | Th/U | 207Pb/235U | 2σ | 208Pb/238U | 2σ | rho | Age (Ma) |
|------------|--------|-------|------|------------|----|------------|----|-----|----------|
| UBV-01     | 2.97   | 5.39  | 0.55 | 2.48712    | 0.0896 | 0.2119 | 0.0027 | 0.355 | 1333.8   |
| UBV-02     | 3.73   | 3.67  | 1.02 | 2.54174    | 0.1523 | 0.2009 | 0.0031 | 0.259 | 1464.2   |
| UBV-03     | 1.57   | 1.68  | 0.94 | 2.51184    | 0.1322 | 0.2141 | 0.0031 | 0.278 | 1421.3   |
| UBV-04     | 2.16   | 3.13  | 0.69 | 2.506      | 0.1319 | 0.2167 | 0.0032 | 0.277 | 1342     |
| UBV-05     | 4.63   | 7.04  | 0.66 | 2.53878    | 0.1149 | 0.2118 | 0.0029 | 0.307 | 1403.5   |
| UBV-07     | 1.41   | 2.29  | 0.62 | 2.53318    | 0.1291 | 0.2192 | 0.0032 | 0.284 | 1329.9   |
| UBV-08     | 4.44   | 9.36  | 0.47 | 2.4279     | 0.1333 | 0.1769 | 0.0028 | 0.267 | 1495.7   |
| UBV-10     | 3.11   | 5.23  | 0.59 | 2.61489    | 0.1054 | 0.2115 | 0.0029 | 0.335 | 1378.4   |
| UBV-11     | 6.07   | 7.16  | 0.85 | 2.51741    | 0.1131 | 0.2129 | 0.0030 | 0.312 | 1373.5   |
| UBV-12     | 11.33  | 7.41  | 1.53 | 2.46729    | 0.1101 | 0.2073 | 0.0029 | 0.314 | 1379.7   |
| UBV-13     | 2.19   | 3.17  | 0.69 | 2.59442    | 0.1152 | 0.2126 | 0.0029 | 0.304 | 1410.7   |
| UBV-14     | 2.70   | 4.38  | 0.62 | 2.56869    | 0.0982 | 0.2152 | 0.0028 | 0.339 | 1373.3   |
| UBV-15     | 5.93   | 7.03  | 0.84 | 2.55049    | 0.1226 | 0.2172 | 0.0030 | 0.289 | 1381     |
| UBV-17     | 7.41   | 10.41 | 0.71 | 2.38887    | 0.0926 | 0.1798 | 0.0024 | 0.337 | 1593.1   |
| UBV-18     | 13.44  | 14.91 | 0.90 | 2.20437    | 0.0684 | 0.1791 | 0.0022 | 0.396 | 1402.4   |
| UBV-19     | 10.41  | 12.40 | 0.84 | 2.58039    | 0.0986 | 0.2011 | 0.0026 | 0.338 | 1465.7   |
| UBV-20     | 4.98   | 7.23  | 0.69 | 2.41636    | 0.0966 | 0.1829 | 0.0024 | 0.330 | 1571.2   |
| UBV-21     | 3.42   | 2.35  | 1.45 | 2.51246    | 0.1883 | 0.2147 | 0.0038 | 0.233 | 1441.6   |
| UBV-22     | 4.25   | 4.43  | 0.96 | 2.29574    | 0.1139 | 0.1833 | 0.0026 | 0.286 | 1395.7   |
| UBV-23     | 3.42   | 5.62  | 0.61 | 2.51595    | 0.2065 | 0.2091 | 0.0038 | 0.223 | 1371.2   |
| UBV-24     | 5.74   | 14.00 | 0.41 | 2.15522    | 0.0801 | 0.1794 | 0.0022 | 0.336 | 1380.2   |
| UBV-25     | 20.21  | 11.32 | 1.79 | 2.04652    | 0.0650 | 0.1590 | 0.0019 | 0.378 | 1386.4   |
| UBV-27     | 16.71  | 23.89 | 0.70 | 2.17854    | 0.0907 | 0.1822 | 0.0024 | 0.311 | 1346.1   |
| UBV-29     | 34.48  | 33.68 | 1.02 | 2.10274    | 0.0678 | 0.1721 | 0.0021 | 0.375 | 1400.8   |
| UBV-30     | 10.94  | 11.12 | 0.98 | 2.25221    | 0.1142 | 0.1828 | 0.0026 | 0.275 | 1370.5   |
| UBV-31     | 29.98  | 40.14 | 0.75 | 2.1384     | 0.0708 | 0.1712 | 0.0021 | 0.367 | 1444     |
| UBV-32     | 11.17  | 16.27 | 0.69 | 2.06116    | 0.0838 | 0.1744 | 0.0023 | 0.317 | 1356.2   |
| UBV-33     | 16.56  | 21.94 | 0.75 | 2.13549    | 0.0722 | 0.1755 | 0.0022 | 0.362 | 1410.4   |
| UBV-34     | 6.19   | 9.48  | 0.65 | 2.14217    | 0.0831 | 0.1703 | 0.0022 | 0.329 | 1436.4   |
| UBV-35     | 3.15   | 4.86  | 0.65 | 2.26149    | 0.0889 | 0.1889 | 0.0024 | 0.326 | 1401.9   |
| UBV-36     | 149.88 | 55.01 | 2.72 | 1.61357    | 0.0579 | 0.1109 | 0.0014 | 0.349 | 1545.2   |

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the Cross, Iddings, Pirsson and Washington norm, the normative Qz–Ab–Or ternary diagram (Johannes & Holtz 1996), the granite’s crystallization pressure varies from 1 to 2 GPa (Fig. 10f).

6.d. Age constraints

The southeastern margin of India has witnessed several extensional as well as subduction events. The first extension- and rift-related Palaeoproterozoic anorogenic magmatism occurred at 1.9 Ga due to mantle diapir or plume. This rifting would have allowed an open ocean to form on the eastern margin of the EDC. The occurrence of 1.85 Ga Kandra and 1.33 Ga Kanigiri ophiolite affinities rocks in the NSB indicates the repeated subduction-related tectonic convergence. The Mesoproterozoic rift-related magmatism occurred in PAkP, resulting in the emplacement of 1321 Ma Elchuru nepheline syenite, 1334 Ma Purimetla gabbro, 1352 Ma Errakonda ferrosyenite and 1356 Ma Uppalapadu nepheline syenite. We find that the plutons emplaced around the Kanigiri area, such as 1284 Ma Kanigiri granite, 1256 ± 6 Ma Punugodu granite (present study) and 1251 Ma Peddcharlapalle gabbro, are relatively younger (~ 80 Ma) than the PAkP alkali rocks. There is a sort of trend of emplacement age of plutons ending towards the south. Sain & Saha (2018) have opined that the 1284 Ma Kanigiri granite intruded the 1334 Ma Kanigiri Ophiolite and thus affirmed their post-tectonic emplacement. In line with the above argument, the studied PG with age 1257 ± 6 Ma has intrusive relationships with the Mesoproterozoic (c. 1334 Ma) KO of oceanic arc supra-subduction affinity. The PG probably emplaced during an extensional tectonic environment along the western margin of the Neoarchaean NSB in the eastern Dharwar Craton.

7. Conclusions

A new age of 1257 ± 6 Ma for the Punugodu granite located in the NSB of Southern India is reported. The geochemical characters of the PG broadly align its nature with the metaluminous, alkalic, within-plate A-type granite. Xenoliths hosted in PG are pre-existing felsic metavolcanic rock fragments. The MME and HME represent distinct batches of mantle-derived mafic–felsic (hybrid) magmas respectively that injected, mingled and undercooled within the felsic host PG magma. The FME appears to be early crystallized cogenetic facies of the PG pluton. Y/Nb ratios (Y/Nb < 1.2) indicate an oceanic island basalt (OIB)-like source for the PG melts formed in an anorogenic within-plate setting. Field observations suggest that PG (c. 1257 Ma) has intrusive relationships with the Mesoproterozoic (c. 1334 Ma) KO of oceanic arc supra-subduction affinity. The PG probably emplaced during an extensional tectonic environment along the western margin of the Neoarchaean NSB in the eastern Dharwar Craton.

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Geochemistry and geochronology of Mesoproterozoic Punugodu granite pluton

Fig. 10. (Colour online) [a] Zr and (b) Nb vs 10 000 × Ga/Al discrimination diagrams of Whalen et al. (1987) showing the A-type nature of the Punugodu granite. [c, d] Chemical classification diagrams of A-type granite (after Eby, 1992): (c) Nb–Y–3Ga ternary diagram and (d) Nb–Y–Zr/4 ternary diagram. (e) Zirconium saturation level (ppm) as a function of cationic ratio $M = ([Na + K + 2Ca]/[Al + Si]$ (Watson & Harrison 1983) showing crystallization temperature of granite. (f) The normative Qtz–Ab–Or ternary diagram shows the pressure of crystallization of PG (Johannes & Holtz 1996).

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