Geochemistry and petrogenesis of Proterozoic granitic rocks from northern margin of the Chotanagpur Gneissic Complex (CGC)

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This study presents the geochemical characteristics of granitic rocks located on the northern margin of Chotanagpur Gneissic Complex (CGC), exposed in parts of Gaya district, Bihar and discusses the possible petrogenetic process and source characteristics. These granites are associated with Barabar Anorthosite Complex and Neo-proterozoic Munger–Rajgir group of rocks. The granitic litho-units identified in the field are grey, pink and porphyritic granites. On the basis of geochemical and petrographic characteristics, the grey and pink granites were grouped together as GPG while the porphyritic granites were named as PG. Both GPG and PG are enriched in SiO₂, K₂O, Na₂O, REE (except Eu), Rb, Ba, HFSE (Nb, Y, Zr), depleted in MgO, CaO, Sr and are characterised by high Fe* values, Ga/Al ratios and high Zr saturation temperatures (GPG_{avg} ~ 861°C and PG_{avg} ~ 835°C). The REE patterns for GPG are moderately fractionated with an average (La/Yb)_N ~ 4.55 and Eu/Eu* ~ 0.58, than PG which are strongly fractionated with an average (La/Yb)_N ~ 31.86 and Eu/Eu* ~ 0.75. These features indicate that the granites have an A-type character. On the basis of geochemical data, we conclude that the granites are probably derived from a predominant crustal source with variable mantle involvement in a post-collisional setting.

1. Introduction

The Proterozoic Eon is marked by the voluminous addition of granitic plutons along the continental arcs and the study of their genesis holds the key to our understanding of the evolution of continental crust (Kemp and Hawkesworth 2003; Frost and Frost 2013). The Chotanagpur Gneissic Complex (CGC) in the eastern Precambrian tract of peninsular India has experienced many episodes of felsic magmatism during the Proterozoic. This is evident from a variety of Proterozoic felsic rocks

exposed in different parts of CGC in the form of gneisses and granites. As per Mahadevan (2002), widespread granitic activity in CGC occurred during 1100–1300 and 1600–1500 Ma.

The northern margin of CGC comprises of Bihar Mica Belt (BMB), the Munger Group and the Rajgir–Gaya sub-basins, showing a basement-cover relationship with the CGC gneisses, though most of the field relationship is obscured by the Quaternary alluvium (figure 1) (Ghose and Mukherjee 2000). The study area at the northern margin of CGC near Gaya, Bihar (figure 1 and 2) exposes

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Figure 1. (a) Outline map of India showing the location of different cratons and mobile belts with respect to Chotanagpur Gneissic Complex (CGC). Abbreviations. SB: Satpura Belt; MB: Mahakoshal Belt; ADBM: Aravalli–Delhi Mobile Belt; NSMB: North Singhbhum Mobile Belt; EGB: Eastern Ghats Belt; SMGC: Shillong Meghalaya Gneissic Complex; SONA: Son Narmada Graben; BKC: Bundelkhand Craton; SC: Singhbhum Craton; BC: Bastar Craton; DC: Dharwar Craton. (b) Geological map of the northern part of CGC and adjoining regions showing the location of the study area in the box marked 'c' (after Chatterjee and Ghose 2011). The tectonic features are marked after Sastri *et al.* (1971), Rao (1973), Valdiya (1976) and GSI (2000).

granitic plutons which are composed of different types of granitic litho-units, related pegmatite and quartz veins (figure 2). Chatterjee and Ghose (2011) reported a crystallisation age of 1697 ± 17 Ma for monazites within the porphyritic granites from Barabar Hills near Gaya which confirms the Proterozoic age of granitic plutons in the region. Wanjari *et al.* (2012) too have reported an Rb–Sr whole rock isochron age of 1337 ± 26 Ma for the granites from Paharpur near Gaya (figure 2). The available geochronological data from other parts of CGC based on Rb–Sr systematics yield similar ages for gneiss from central (1717 ± 102 Ma,

Mallik *et al.* 1991) and granites from the western (1741 ± 65 Ma, Ray Barman and Bishui 1994), northern (1590 ± 30 Ma, Pandey *et al.* 1986) and eastern (1599 ± 33 Ma, Mallik *et al.* 1991) parts, which marks the oldest time span of felsic magmatism. However, the last significant granitic and pegmatitic activity marked by Rb–Sr dates is between 950 and 800 Ma (Mahadevan 2002). In this paper, we present a new whole rock major and trace elemental data for granites from Gaya. The dataset allows us to characterise these granitoids and help us understand their petrogenesis and put constraints on tectonic settings.

Figure 2. Detailed geological map of the study area showing various granitoid hillocks north of CGC in close proximity to Rajgir fold belt (modified after parts of Quadrangle Geological Map of 72 H and 72 D). Locations of the samples presented in the study have been marked in the map with solid star (*).

2. Geological background

Chotanagpur Gneissic Complex (CGC), is an ENE–WSW trending high-grade terrain in the eastern part of Central Indian Tectonic Zone (CITZ) and covers an area of about 80,000 km² (figure 1). It is bordered by the Singhbhum mobile belt in the south, the Quaternary Gangetic alluvium in the north and by the Rajmahal basalt in the northeast. In the west, CGC is separated from CITZ by younger Gondwana sediments. The terrain consists of mostly gneisses and migmatites with numerous metasedimentary enclaves of varying metamorphic grade which are in turn intruded by metabasic, anorthositic and granitic plutons, along with widespread but narrow belts of supra-crustal enclaves, discontinuous boudin-type granulite belts and major parallel shear zones with a strike extension of about 500 km and a width of 200 km in E–W direction. Younger mafic, ultramafic and alkaline (sodic and ultrapotassic) intrusives of early Tertiary age are also present (Dunn 1929; Ghose 1983, 1992; Mahadevan 2002; Mazumdar 1988; Sarkar 1988). The rocks of CGC are complexly deformed and metamorphosed showing varying degrees of metamorphism, exhibiting mainly amphibolite facies but ranges from greenschist (mostly in SE part of CGC) to granulite facies (central and eastern parts of CGC) (Bhattacharya 1976; Banerji 1991; Mahadevan 2002; Sharma 2009). Detailed structural studies

have identified three phases of deformation in the CGC, giving rise to distinct fold patterns and related linear fabric (Chatterjee *et al.* 2008 and the references therein).

Based on the available geochronological data on the rocks of CGC, the terrain appeared to have experienced three tectono-thermal events during the Proterozoic. The first tectono-thermal event corresponds to the age group of 1.7–1.6 Ga while the second 1.2–1.0 Ga and third corresponds to 0.9 Ga (Chatterjee *et al.* 2008; Chatterjee and Ghose 2011; Karmakar *et al.* 2011; Sanyal and Sengupta 2012). In the wake of the complexity attained by the rocks of this terrain due to polyphase deformation, metamorphism, magmatism, metasomatism, lack of structural, geochemical, petrological and precise geochronological data, it is hard to ascertain the geological relationship between different litho-units and comment on the sequence of geological events in the region.

The study area, around Gaya, to the north of CGC, predominantly comprises of granitic plutons situated between the ENE and WSW trending Bihar Mica Belt (BMB) in the south and Barabar Anorthosite Complex in the north, whereas on the eastern margin, the Munger Group and the Rajgir–Gaya Basin are situated (figure 1). The Munger Group and the Rajgir–Gaya Basin comprise of folded phyllite-quartzite sequences metamorphosed into greenschist facies with minor intrusive mafic–ultramafic rocks and granitic bodies (Das 1967;

Mazumdar 1988). The age and correlation of these rocks with other rocks of the same area are uncertain (Chatterjee and Ghose 2011). However, Das (1967) has correlated the Munger and Rajgir–Gaya sequences with Son–Narmada valley of central India, based on lithological and structural similarities. In the study area, out of three major sub-vertical NE–SW trending faults present (figure 1), the NW-dipping Monghyr–Saharsa Ridge Fault and SE-dipping East Patna Fault taken together define a graben which probably represents a north-eastward extension of the Son–Narmada Graben of central India. Therefore, the granitic plutons along with the schist-phyllite-quartzite country rocks in the study area, lying on the northwestern side of the Monghyr–Saharsa Ridge Fault of the Rajgir–Gaya Basin may be correlated with the Mahakoshal Group of the Son–Narmada Graben (Chatterjee and Ghose 2011 and the references therein). Further, Chatterjee and Ghose (2011) have correlated these country rocks hosting the porphyritic granites around Gaya with Mahakoshal granitoids and supracrustals on the basis of 1697 ± 17 Ma monazite crystallisation age, and the absence of <1.6 Ga metamorphic imprint in the porphyritic granites.

2.1 Field relationship

The studied granitic plutons lie in parts of the Survey of India Toposheets No. 72 D/13, 72 D/14 and 72 H/1 in Gaya district, Bihar (figure 2) and their exposures form circular to elliptical residual dome-like structures or inselbergs (figure 3a). Three types of granitic litho-units identified in the field on the basis of colour and texture are: (1) grey granite, (2) pink granite and (3) porphyritic granite. As the area is mostly covered with alluvium, the field relationship between litho-units are obscured and good exposures are found in pits of numerous quarries. The grey granite comprises two variants, the first one is leuco- to mesocratic, coarse-grained granites with an inequigranular texture. The second one is more widespread, leucocratic, medium-to-fine grained, equigranular granites which is seen intruding the first one (figure 3b). These two variants of grey granite are undifferentiated in the geological map and are observed around Gandhar, Kalauna, Kapildhara, Paharpur, Pahara and Pretshila (figure 2). These variants of granite are also seen intruding the anorthosites near Sapneri and Lachhubigha and volcano-sedimentary sequences near Bathani. Porphyritic granite occur as massive bodies, but at places, they are weakly foliated, sheared and are known to occur around Bajubigha, Bara and Matua (figure 2). At places, these granites are seen intruded by fine-grained grey granites

(figure 3c), and are characterised by sub-rounded to tabular shaped phenocrysts of feldspars typically ranging in size from 2–4 cm in diameter (figure 3e), but sometimes they can measure up to 5–6 cm (figure 3d). These phenocrysts constitute about 30–40% of the total mass and are randomly arranged, but at places, magmatic foliation aligned parallel to NE regional foliation trend is recorded. Occasionally rapakivi-like texture is observed in the feldspar phenocrysts. Pink granites are generally coarse to medium-grained, equigranular and homogeneous in nature (figure 3f) and are known to occur around Korma and Gere (figure 2). These granites display a clear inter-fingering contact with the rhyolite near Gere (figure 2). The Rb–Sr whole-rock dating of these rhyolites yields a crystallisation age of 1044 ± 35 Ma (Wanjari *et al.* 2012). Mafic magmatic enclaves (MME's) of variable sizes and angular to sub-rounded shapes are commonly seen in all the granitic litho-units. All these granitic plutons are interspersed with fine-grained aplitic or coarse-grained pegmatitic veins at places.

Wanjari *et al.* (2012) have reported pillow basalts and associated pyroclasts in the vicinity of Gaya along with rhyolite from Gere and Mirzapur and even suggested a back-arc basin like tectonic setting for the pillow basalts and mafic pyroclasts. Reports of occurrence of Bathani volcano-sedimentary sequences (BVS), northeast to our study area in juxtaposition to the Munger–Rajgir metasedimentary fold belt also adds emphasis to this study (Ahmad and Wanjari 2009; Saikia *et al.* 2014). This volcano-sedimentary sequence comprised of garnet-mica schist, banded iron formation, chert bands, volcanic tuff, associated mafic pyroclasts and pillow basalts. Saikia *et al.* (2014) proposed a multi-stage and multi-source magmatism for BVS in an island-arc tectonic setting. It is thus imperative to envisage a viable geological model for the evolution of granitic rocks around Gaya and their interrelationships. Therefore, constraining the geochemical signatures of these granitic rocks will help us in understanding the geodynamic evolution of this area.

3. Petrography

3.1 Grey granites

Mineralogically, both types of grey granites are similar and shows holocrystalline, seriate to hypidiomorphic texture, and essentially consists of Qtz (34 vol.%) + Pl (28 vol.%) + Kfs (24 vol.%) + Hbl (9 vol.%) + Bt (4 vol.%) (mineral symbols after Siivola and Schmid 2007) (figure 4a). Accessory minerals (<1 vol.%) include primary Ttn + Ap + Zrn ± Aln + Op, and secondary Ep + Chl + Ser ± Ms. Quartz is present as strained grains and includes

incipient sub-grains. Plagioclase, mostly unzoned and poikilitic, shows moderate alteration to sericitization and saussuritization in some places. K-feldspar is mostly microcline and shows variable perthitic development with well-developed veins and flame to patchy perthites (figure 4b). Myrmekitic

texture and micrographic texture is common. Euhedral to subhedral hornblende poikilitically encloses biotite, zircon, apatite and opaques, and coexist mostly in association with green to straw brown biotite (figure 4c). Biotite, at places, shows alteration to chlorite (figure 4d).

Figure 3. Field photographs of various granitic litho-units showing (a) exposure of granitic pluton near Bara, (b and c) cross-cut relationship of fine-grained grey granite which intrudes both the coarse-grained grey granite and porphyritic granite. The inset in (c) shows close-up contact of fine grained grey granite and porphyritic granite, (d) phenocrysts of K-feldspar in porphyritic granite, (e) mafic microgranular enclave (MME) in porphyritic granite, and (f) outcrop of pink granite.

Figure 4. Photomicrographs of granitic litho-units showing (a) hypidiomorphic texture, minerals present are plagioclase, K-feldspar, quartz, hornblende and biotite (sample B-15), (b) perthitic intergrowth in K-feldspar with plagioclase and quartz grains (sample B-11), (c) hornblende and biotite grains in the matrix and associated epidote needles (sample B-12), (d) chlorite as alteration product of biotite is seen along with sericitised plagioclase and quartz grains (sample B-23), (e) myrmekite development at the boundary of K-feldspar (sample B-8), and (f) granophyric texture (sample B-10). (Qtz: quartz; Kfs: K-feldspar; Pl: plagioclase; Bt: biotite; Hbl: hornblende; Ep: epidote; Chl: chlorite).

3.2 Pink granites

The pink granites show hypidiomorphic texture, composed of Qtz (36 vol.%), Kfs (29 vol.%), Pl (25 vol.%), Bt (7 vol.%) and Hbl (2 vol.%). Accessory

minerals (<1 vol.%) include only Zrn (commonly in biotite forming pleochroic halos)+Ap+Op±Aln. K-feldspar is more common than plagioclase, poikilitic and mainly shows simple twinning, but at places tartan twinning typical of microcline is

also noticed. Exsolution and intergrowth textures like perthitic and granophyric are very common (figure 4f). These granites show microstructural deformation features, viz., strained quartz grains, kinked mica and deformation twins in plagioclase.

3.3 Porphyritic granites

The porphyritic granites are composed of Kfs (25 vol.%) and Pl (10 vol.%) phenocrysts set in a coarse grained matrix consisting of Qtz (26 vol.%), Pl (19 vol.%), Kfs (10 vol.%), Bt (8 vol.%) and minor Hbl (<1 vol.%). They also contain accessory (<1 vol.%) Zrn+Aln+Ap+Ttn+Op minerals. Phenocrysts of K-feldspar are mostly microcline and at places present with perthitic exsolution lamellae. Myrmekitic intergrowth is common along the grain boundaries of K-feldspar (figure 4e), and at places micrographic texture is present. The effects of shearing and deformation are evidenced by the presence of fine-grained recrystallised quartz, deformation twinning in feldspar, kinked mica, undulose extinction in quartz and development of a weak foliation defined by alignment of biotite.

4. Analytical methods

In this study, 32 samples (approx. 5–7 kg each) were collected but a total of 27 samples (15 grey granites, 5 pink granites and 7 porphyritic granites) with least alteration and deformation were analysed for major oxides, trace elements and rare earth elements (table 1). The average chemical index of alteration (CIA) value of all samples is 53, which lies within the accepted 45–55 range for unweathered granites (Nesbitt and Young 1982). Loss on ignition (LOI) was determined by calcining the samples for 2 hours at 900°C in an electric furnace. Whole rock powders were analysed for major and most trace elements using WD-XRF (PANalytical AXIOUS PW4400/40 spectrometer) at the Department of Geology, University of Delhi, India. The accuracy of the analyses for the major elements is better than 1% for SiO₂ and 2% for other major elements, 2–5% for minor elements and better than 10% for trace elements. Some trace and rare earth elements were analysed by ICP-MS (Perkin-Elmer Sciex ELAN DRC 6000) at the Institute Instrumentation Centre (IIC), Indian Institute of Technology Roorkee, India. Protocols for sample digestion and preparation of solutions for ICP-MS were that of Balaram *et al.* (1996). The international rock standards (USGS) used for calibrations for both WD-XRF and ICP-MS are JG-2 and DG-H. The precision of the data was expressed as the residual standard deviation (RSD), which in general for all the trace elements is much less than

4%. Individually, they are less than 2.5% for alkali and alkaline earth metals, whereas it is less than 2% for transition elements except Sc and Hf and within 3% for LREE and 4% for HREE (Bhattacharya *et al.* 2012).

5. Geochemical characteristics

On the basis of geochemical compositions and petrographic characteristics, grey granites (both, coarse and fine-grained variants) and pink granites have been classified together and herein named as GPG (20 samples) and the porphyritic granites have been named as PG (7 samples). Major and trace element data for all granites are listed in table 1.

5.1 Major and trace elements

The major and trace element distinctions between GPG and PG are subtle, but the real notable differences come in REE patterns. GPG and PG samples show similar abundances with high SiO₂ (GPG_{avg} = 72.75 wt.%; PG_{avg} = 69 wt.%), Fe₂O₃^t (GPG_{avg} = 3.75 wt.%; PG_{avg} = 3.51 wt.%), Al₂O₃, Na₂O, K₂O, low MgO and CaO. However, on an average PG have higher Al₂O₃, TiO₂, MgO and P₂O₅ content than GPG. Collectively, samples of both GPG and PG show high average contents of Rb (~228 ppm), Ba (~529 ppm), high field strength elements (HFSE) (Nb~21 ppm, Zr~307 ppm, Y~48 ppm, Th~25 ppm and U~3 ppm) and depletion in Sr (~56 ppm). All samples of GPG and PG have normative quartz, corundum and hypersthene. In the normative QAP classification diagram (figure 5a), all samples plot in the monzogranite field except sample B-23 which lies in the granodiorite field. On the other hand, on the Ab–An–Or diagram of O'Connor (1965) modified after Barker (1979) (figure not shown), all samples plot in granite field. The samples of GPG are slightly metaluminous to peraluminous (A/CNK = 0.97–1.36) while those of PG are peraluminous (A/CNK = 1.07–1.21) (figure 5b). According to the classification of Frost *et al.* (2001), all samples belong to the ferroan granite on Fe*[FeO^t/(FeO^t+MgO)] *vs.* SiO₂ diagram (figure 5c), whereas on the MALI (Na₂O+K₂O–CaO) *vs.* SiO₂ (figure 5d), samples of GPG mostly plot in the calc-alkalic and those of PG plot in alkalic-calcic fields. These observations are consistent with the chemical characteristics of worldwide A-type granites. Although the granites have high Fe*, interestingly they are K-rich and their average K₂O/Na₂O~1.64 ratios are quite high. This indicates that the granites in the study area are roughly similar to the peraluminous A-type granites.

Table 1. *Representative whole rock compositions of granites from Gaya.*

Sample	B-02	B-05	B-13	B-14	B-15	B-19	B-21	B-22	B-23
Rock type	FGG	FGG	FGG	FGG	FGG	FGG	FGG	FGG	FGG
Major oxides (wt.%)									
SiO ₂	70.98	76.03	71.93	73.07	68.02	77.06	75.60	74.97	64.62
Al ₂ O ₃	13.18	13.83	13.86	13.57	13.75	14.51	13.90	14.14	15.61
Na ₂ O	2.95	3.30	2.69	2.88	3.15	3.17	2.90	2.98	3.59
MgO	0.16	0.03	0.05	0.11	0.16	0.06	0.07	0.06	1.03
MnO	0.09	0.05	0.08	0.09	0.12	0.05	0.04	0.04	0.04
Fe ₂ O ₃ ^t	5.08	2.61	4.67	4.63	5.88	2.74	2.45	2.41	6.07
K ₂ O	4.62	4.66	4.81	4.67	4.62	5.03	5.46	5.34	2.53
CaO	1.72	0.96	1.69	1.43	2.19	1.06	0.99	1.02	2.07
TiO ₂	0.34	0.20	0.35	0.31	0.40	0.21	0.20	0.20	0.52
P ₂ O ₅	0.06	0.02	0.03	0.03	0.06	0.03	0.03	0.03	0.05
LOI	0.83	0.51	0.76	0.72	0.68	0.55	0.52	0.45	1.95
Total	100.00	102.20	100.92	101.50	99.03	104.47	102.15	101.63	98.06
CIA									
CIA	50	53	52	52	49	54	53	53	56
Fe*									
Fe*	0.97	0.99	0.99	0.97	0.97	0.97	0.97	0.97	0.84
MALI									
MALI	5.86	7.00	5.82	6.11	5.58	7.15	7.36	7.30	4.05
Trace elements (ppm)									
Rb	204	240	178	209	241	243	226	239	170
Ba	750	649	678	623	563	522	516	737	561
Th	8	16	21	23	21	19	19	8	25
U	3	2	2	4	5	2	2	1	9
Nb	20	18	16	23	29	18	17	18	25
Pb	29	24	24	27	24	27	22	22	19
Sr	87	34	51	61	70	35	36	41	66
Zr	294	328	315	360	404	344	324	300	509
Hf	7	6	6	6	6	6	6	6	3
Y	52	54	46	57	63	61	57	52	70
Ni	6	21	24	15	5	10	24	15	19
Cr	651	733	522	779	580	726	557	518	419
V	11	5	10	9	16	8	6	2	34
Co	8	4	8	5	4	6	6	5	0
Sc	21	2	1	12	15	7	4	9	0
Cu	18	7	36	6	10	7	3	1	0
Ga	22	22	23	23	23	23	23	22	25
Zn	94	82	78	108	142	85	62	78	27
La	30.70	26.21	20.56	27.24	24.82	24.58	24.22	22.93	33.63
Ce	69.27	58.80	46.55	61.06	56.02	54.49	55.31	52.54	75.64
Nd	39.74	33.54	26.99	35.31	32.61	30.93	32.10	30.68	43.77
Sm	9.02	7.53	6.24	8.13	7.60	6.88	7.43	7.19	10.09
Eu	2.27	1.49	1.45	1.93	2.03	1.29	1.38	1.64	2.07
Gd	10.07	8.38	7.17	9.43	8.82	7.92	8.39	8.15	11.13
Tb	1.55	1.27	1.11	1.47	1.37	1.21	1.29	1.23	1.69
Dy	9.89	7.82	7.04	9.49	9.11	7.69	8.09	7.53	10.38
Ho	1.95	1.51	1.39	1.87	1.85	1.51	1.59	1.47	2.02
Er	5.01	3.79	3.56	4.80	4.94	3.84	4.04	3.70	5.07
Yb	5.31	3.65	3.64	5.02	5.92	3.86	4.04	3.60	5.69
Lu	0.83	0.54	0.55	0.76	0.92	0.58	0.62	0.55	0.82
Eu/Eu*									
Eu/Eu*	0.73	0.57	0.66	0.67	0.76	0.53	0.53	0.65	0.60
(La/Lu) _N									
(La/Lu) _N	3.86	5.02	3.85	3.70	2.79	4.44	4.07	4.36	4.25
T _{Zr} (°C)									
T _{Zr} (°C)	841.61	868.71	857.10	872.21	862.90	874.62	865.26	858.66	914.99

Table 1. (Continued.)

Sample	B-11	B-12	B-24	B-25	B-31	B-32	B-03	B-10	B-27
Rock type	CGG	CGG	CGG	CGG	CGG	CGG	PG	PG	PG
Major oxides (wt.%)									
SiO ₂	74.97	65.45	70.79	72.40	75.18	72.40	76.19	73.50	73.98
Al ₂ O ₃	13.89	13.98	13.88	13.99	13.75	14.11	13.45	14.07	14.39
Na ₂ O	3.03	2.95	2.87	2.28	3.44	2.88	3.49	3.14	3.25
MgO	0.17	0.27	0.72	0.75	0.17	0.33	0.35	0.13	0.20
MnO	0.05	0.14	0.01	0.03	0.02	0.03	0.01	0.04	0.06
Fe ₂ O ₃ ^t	3.27	7.82	3.87	3.84	2.11	2.81	2.78	3.01	3.18
K ₂ O	4.94	3.57	4.67	5.45	4.32	5.20	4.71	5.14	4.94
CaO	1.21	2.68	0.78	0.37	1.04	0.94	0.49	1.08	1.22
TiO ₂	0.24	0.54	0.26	0.26	0.14	0.23	0.21	0.22	0.20
P ₂ O ₅	0.03	0.09	0.03	0.03	0.02	0.04	0.02	0.03	0.02
LOI	0.64	0.87	1.08	1.09	1.00	0.79	0.40	0.82	1.12
Total	102.45	98.35	98.95	100.49	101.19	99.77	102.10	101.17	102.54
CIA									
CIA	53	51	55	58	53	54	53	53	53
Fe*									
Fe*	0.94	0.96	0.83	0.82	0.92	0.89	0.88	0.96	0.93
MALI									
MALI	6.76	3.84	6.77	7.37	6.72	7.15	7.71	7.19	6.96
Trace elements (ppm)									
Rb	238	174	210	166	185	221	102	279	217
Ba	706	456	386	540	143	272	412	710	662
Th	24	19	21	24	42	40	22	20	25
U	4	3	3	2	8	7	1	4	4
Nb	22	26	23	20	17	16	16	23	23
Pb	26	23	10	12	35	33	12	23	29
Sr	48	90	37	27	23	36	31	34	37
Zr	318	393	315	335	152	169	391	301	350
Hf	7	5	6	7	7	10	6	7	10
Y	58	44	47	32	77	73	31	66	62
Ni	20	14	17	22	7	24	21	21	14
Cr	604	386	562	407	881	760	828	677	486
V	15	20	12	15	6	19	11	10	8
Co	4	4	6	5	4	8	4	5	8
Sc	11	9	9	3	2	0	9	23	14
Cu	2	39	0	44	1	5	10	15	1
Ga	21	23	21	18	21	20	23	23	23
Zn	85	132	17	13	24	39	13	58	89
La	30.46	21.79	28.81	26.06	24.99	22.63	34.83	27.50	27.66
Ce	67.04	50.65	62.61	56.64	51.02	47.04	74.97	60.70	61.27
Nd	37.40	30.25	33.84	30.21	25.83	23.66	39.19	34.22	34.83
Sm	8.04	7.37	6.80	5.89	4.41	3.97	7.28	7.51	7.77
Eu	1.67	2.31	1.35	1.70	0.24	0.41	1.49	1.50	1.59
Gd	9.14	8.38	7.58	6.58	5.44	4.82	8.20	8.47	8.87
Tb	1.35	1.33	1.06	0.82	0.79	0.71	1.11	1.30	1.37
Dy	8.54	8.40	6.16	4.44	5.16	4.69	6.37	8.44	8.89
Ho	1.69	1.67	1.21	0.85	1.10	1.00	1.23	1.68	1.79
Er	4.32	4.32	3.08	2.08	3.11	2.81	3.09	4.35	4.70
Yb	4.50	4.71	3.19	2.15	3.45	3.33	3.13	4.64	5.04
Lu	0.69	0.74	0.51	0.36	0.52	0.49	0.49	0.71	0.77
Eu/Eu*									
Eu/Eu*	0.59	0.90	0.57	0.83	0.15	0.28	0.59	0.57	0.58
(La/Lu) _N									
(La/Lu) _N	4.59	3.07	5.90	7.53	4.95	4.81	7.32	4.04	3.74
T _{Zr} (°C)									
T _{Zr} (°C)	862.32	864.98	871.01	887.22	795.69	806.22	888.20	855.64	870.95

Table 1. (*Continued.*)

Sample	B-28	B-30	B-06	B-07	B-08	B-09	B-16	B-17	B-18
Rock type	PG	PG	PG*	PG*	PG*	PG*	PG*	PG*	PG*
Major oxides (wt.%)									
SiO ₂	70.02	77.85	69.38	77.09	62.67	65.79	68.37	69.95	69.78
Al ₂ O ₃	14.57	14.13	15.65	13.83	15.30	14.96	14.68	15.37	15.70
Na ₂ O	3.20	3.09	2.84	2.46	2.76	2.95	2.94	2.42	2.73
MgO	0.71	0.11	0.64	0.14	1.06	1.02	0.71	0.29	0.51
MnO	0.03	0.05	0.03	0.02	0.07	0.07	0.05	0.02	0.03
Fe ₂ O ₃ ^t	4.31	1.55	3.61	1.36	5.56	5.20	4.10	1.78	2.96
K ₂ O	3.65	5.57	5.12	5.45	4.83	4.97	4.46	6.85	5.93
CaO	1.97	0.58	1.90	0.80	2.14	2.09	2.03	1.03	1.39
TiO ₂	0.51	0.14	0.43	0.16	0.71	0.66	0.50	0.18	0.31
P ₂ O ₅	0.08	0.02	0.13	0.03	0.21	0.21	0.14	0.11	0.11
LOI	1.34	1.09	1.06	0.53	0.83	0.75	0.88	0.59	0.77
Total	100.41	104.16	100.78	101.86	96.14	98.67	98.85	98.59	100.23
CIA	53	54	53	55	53	52	52	54	54
Fe*	0.84	0.93	0.83	0.90	0.82	0.82	0.84	0.85	0.84
MALI	4.88	8.08	6.06	7.10	5.45	5.83	5.36	8.24	7.27
Trace elements (ppm)									
Rb	183	205	227	234	252	218	229	278	259
Ba	435	321	583	256	615	902	318	477	610
Th	13	33	35	11	35	11	21	6	31
U	4	7	2	2	1	2	2	1	3
Nb	22	28	17	8	30	27	24	13	21
Pb	20	21	27	21	15	18	20	21	32
Sr	65	21	113	75	89	98	69	72	90
Zr	378	198	335	145	351	243	293	186	253
Hf	5	7	8	5	4	3	5	6	7
Y	44	65	19	19	42	33	28	29	24
Ni	18	15	14	7	4	14	8	4	10
Cr	590	1251	608	752	420	444	576	555	530
V	31	9	30	6	46	50	33	17	19
Co	8	5	9	6	10	9	7	9	4
Sc	4	10	0	6	13	11	12	13	12
Cu	20	3	23	1	11	11	6	0	0
Ga	21	22	23	19	23	20	24	21	20
Zn	37	9	55	17	75	71	58	26	49
La	22.85	28.24	44.79	34.24	54.53	18.69	30.06	12.53	31.87
Ce	48.88	61.98	93.52	68.01	111.04	41.59	61.11	27.05	66.86
Nd	26.52	34.37	44.15	30.78	53.69	22.74	28.97	14.04	32.61
Sm	5.38	7.30	6.02	3.51	7.97	4.69	4.01	2.56	4.98
Eu	1.18	0.60	1.26	1.06	1.31	1.39	0.88	0.89	1.19
Gd	6.45	8.66	6.10	3.87	8.71	5.05	4.72	2.60	4.92
Tb	1.00	1.35	0.48	0.32	1.00	0.69	0.50	0.29	0.42
Dy	6.50	8.77	1.46	1.19	4.91	3.90	2.30	1.46	1.45
Ho	1.31	1.77	0.28	0.23	0.90	0.75	0.43	0.28	0.26
Er	3.42	4.65	0.70	0.57	2.08	1.87	1.04	0.67	0.61
Yb	3.62	5.08	0.52	0.49	1.42	1.85	0.86	0.64	0.46
Lu	0.53	0.76	0.10	0.08	0.21	0.28	0.14	0.10	0.08
Eu/Eu*	0.61	0.23	0.63	0.88	0.48	0.87	0.62	1.05	0.74
(La/Lu) _N	4.45	3.86	44.92	44.42	27.61	6.82	22.13	12.94	40.34
T _{Zr} (°C)	877.70	823.06	864.19	799.69	859.81	821.12	846.88	811.75	839.31

FGG: Fine grained grey granite; CGG: Coarse grained grey granite; PG: Pink granite; PG*: Porphyritic granite.

Figure 5. (a) Q–A–P normative plot after Le Maitre *et al.* (2002) showing granitic composition for the samples, (b) ACNK (molar $\text{Al}_2\text{O}_3/(\text{CaO}+\text{Na}_2\text{O}+\text{K}_2\text{O})$) vs. A/NK (molar $\text{Al}_2\text{O}_3/(\text{Na}_2\text{O}+\text{K}_2\text{O})$) diagram (field after Maniar and Piccoli 1989) showing chemical composition of GPG and PG in terms of alumina saturation, (c) $\text{Fe}^*[\text{FeO}^t/(\text{FeO}^t+\text{MgO})]$ vs. SiO_2 , and (d) MLI-index ($\text{Na}_2\text{O}+\text{K}_2\text{O}-\text{CaO}$) vs. SiO_2 diagrams after Frost *et al.* (2001).

Major elements of GPG and PG form well-defined clusters and trends in Harker bivariate plots (figure 6), where TiO_2 , P_2O_5 , CaO , Fe_2O_3^t and Al_2O_3 concentrations decrease monotonously with increasing amount of SiO_2 . The variation of Na_2O however, is independent of the SiO_2 , and the K_2O displays a weak positive correlation with SiO_2 . The overall pattern of decreasing TiO_2 , Fe_2O_3^t , MgO and CaO with increasing SiO_2 is characteristic of fractionating granitic systems. The variation patterns of selected trace elements with respect to each other and SiO_2 show simple fractionation trends (figure 7). Ba, Sr and Zr generally decrease with increasing SiO_2 for all samples of GPG and PG, whereas Rb shows little correlation with increasing silica. Rb and Ba are more enriched relative to Sr, and overall Rb/Sr ratios are high but comparatively higher in GPG (1.92–9.84) than PG (2.02–3.87). The major and trace elements along with inter-elemental variation diagrams suggest a key role of fractional crystallisation processes during petrogenesis of these granites.

5.2 Trace and Rare earth element patterns

Chondrite normalised REE patterns reveal a distinction between GPG and PG (figure 8a and c), where samples of GPG show similar patterns with a lesser degree of REE fractionation ($(\text{La}/\text{Yb})_N = 2.85\text{--}8.24$) than PG, which shows a higher degree of fractionation ($(\text{La}/\text{Yb})_N = 6.88\text{--}58.12$). The GPG has high ΣREE (an average 153.85 ppm) content, and shows moderate LREE enrichment ($(\text{La}/\text{Sm})_N = 1.85\text{--}3.56$) with enriched but relatively flat HREE ($(\text{Gd}/\text{Yb})_N = 1.17\text{--}2.48$) patterns and distinct negative Eu anomaly ($\text{Eu}/\text{Eu}^* = 0.15\text{--}0.9$) (figure 8a). PG, on the other hand, has relatively low ΣREE (an average 148.40 ppm) contents and exhibits strongly fractionated REE patterns with enriched LREE ($(\text{La}/\text{Sm})_N = 2.49\text{--}6.08$) and depleted HREE ($(\text{Gd}/\text{Yb})_N = 2.21\text{--}9.43$) patterns having concave upward shape, and show weak negative to moderate positive Eu anomaly ($\text{Eu}/\text{Eu}^* = 0.48\text{--}1.05$) (figure 8c). On the primitive mantle normalised multi-element spidergrams

Figure 6. Harker variation diagrams of the major element oxides with SiO_2 for the studied granites.

(figure 8b and d), samples of GPG and PG show similar patterns with more or less same degree of elemental enrichment. They show characteristic features common to A-type granites, viz., distinctive depletion of LILE's like Ba, Sr and HFSE's like Nb, P and Ti relative to other trace elements.

5.3 Classification

All samples of GPG and PG plot in the field of A-type granites in the Fe^* vs. SiO_2 diagram of Frost *et al.* (2001) (figure 5c). Also, Whalen *et al.* (1987) have proposed discrimination diagrams to distinguish A-type granites from the other variants like I-, S- and M-type granites. The variation of the major oxide ratios and trace elements like $(\text{K}_2\text{O} + \text{Na}_2\text{O})/\text{CaO}$, FeO^t/MgO , Zr and Nb with Ga/Al (figure 9a, b, d and e), show that all the samples clearly correspond to A-type granites. However, some less evolved samples lie at the boundary between the A-type and (I, S and M)-type fields in the diagrams. In the $(\text{K}_2\text{O} + \text{Na}_2\text{O})/\text{CaO}$ and FeO^t/MgO vs. $\text{Zr} + \text{Nb} + \text{Ce} + \text{Y}$ diagrams to discriminate fractionated granites from A-type

granites (figure 9c and f), the samples show a consistent A-type character. Moreover, some samples from both GPG and PG have a compositional overlap between A-type and fractionated granites (FG). In Rb vs. $(\text{Y} + \text{Nb})$ tectonic discrimination diagram of Pearce *et al.* (1984) (figure 9g), the majority of GPG samples lie in the within-plate granite field whereas the PG samples straddle at the triple junction boundary of the within-plate, volcanic arc and syn-collisional granites. However, in Y vs. Nb (figure 9h) majority of samples are concentrated predominantly in the within-plate granite field while some samples of GPG fall in the field of volcanic arc plus syn-collisional granites. This also provides support for an A-type character as A-type is considered to be equivalent to within-plate granites (Whalen *et al.* 1987; Eby 1992).

6. Discussion

6.1 Estimated temperature of magma

Zircon saturation temperatures (T_{Zr}) calculated from bulk rock compositions provide a simple and

Figure 7. Binary diagrams of (a) SiO_2 vs. Rb, (b) SiO_2 vs. Ba, (c) Sr vs. Ba, (d) SiO_2 vs. Sr, (e) SiO_2 vs. Rb/Sr, (f) CaO vs. Sr, (g) SiO_2 vs. Zr, (h) MgO vs. Y and (i) SiO_2 vs. Eu/Eu^* showing fractionation of feldspar, hornblende, zircon for the studied granites. The vectors in Sr vs. Ba plot are taken from Huang *et al.* (2012).

robust means of estimating magma temperatures (Watson and Harrison 1983; Miller *et al.* 2003). Watson and Harrison (1983) have established experimentally that the partition coefficient of zircon $D_{\text{Zr}}^{\text{zircon/melt}}$ is a function of the parameter $M[(\text{Na} + \text{K} + 2\text{Ca})/(\text{Al} * \text{Si})]$, all in cation fraction] and temperature. Therefore, using this formula, the Zr thermometry yields the temperature at which the granites of study area formed, i.e., an average T_{Zr} ($^{\circ}\text{C}$) ~ 861 for GPG and 835 for PG (table 1) (figure 9i). However, Miller *et al.* (2003) showed that the average T_{Zr} for inheritance-rich granitoids are 766°C and inheritance-poor granitoids are 837°C because zirconium concentrations in inheritance-rich granitoids are uniformly lower than those of inheritance-poor intrusions. Consequently, calculated T_{Zr} values are also lower for inheritance-rich granitoids. Therefore, the rocks in the study area are hot inheritance-poor granitoids, i.e., undersaturated in zircon at the source, with $T_{\text{Zr}} > 800^{\circ}\text{C}$ and the T_{Zr} suggests a minimum initial magma temperature at the source.

6.2 A-type granites: Reduced or oxidised nature?

The granites in the study area have most of the petrological and geochemical characteristics associated with the definition of A-type granites (Loiselle and Wones 1979; Whalen *et al.* 1987; Eby 1990; Frost *et al.* 2001). They are amphibole-bearing and the bulk composition is characterised by low CaO, Ba and Sr, high $\text{Na}_2\text{O} + \text{K}_2\text{O}$ and $\text{FeO}^t/(\text{FeO}^t + \text{MgO})$, high REE (except Eu) and HFSE (Zr, Y and Nb) values. They also have very high Zr saturation temperatures ($799\text{--}915^{\circ}\text{C}$) and lack of significant inherited zircon also support to an A-type character (Creaser *et al.* 1991). Overall, the granites are very similar in composition and mineralogy to the peraluminous A-type granites from Australia (Turner *et al.* 1992; King *et al.* 1997).

Dall'Agnol and de Oliveira (2007) have proposed a new classification scheme to separate calc-alkaline granites from A-type granites and oxidised A-type granites from reduced A-type granites. In the $\text{CaO}/(\text{FeO}^t + \text{MgO} + \text{TiO}_2)$ vs. $\text{CaO} + \text{Al}_2\text{O}_3$

Figure 8. Chondrite normalised rare earth elements (REE) patterns (a and c) (normalisation value after McDonough and Sun 1995) and primitive mantle normalised trace element patterns (b and d) (normalisation value after Sun and McDonough 1989).

and $\text{CaO}/(\text{FeO}^t + \text{MgO} + \text{TiO}_2)$ *vs.* Al_2O_3 diagrams (figure 10a and b), the granites fall mostly in the A-type granite field and they differ clearly from the calc-alkaline granites. In the $\text{FeO}^t/(\text{FeO}^t + \text{MgO})$ *vs.* Al_2O_3 and $\text{FeO}^t/(\text{FeO}^t + \text{MgO})$ *vs.* $\text{Al}_2\text{O}_3/(\text{K}_2\text{O}/\text{Na}_2\text{O})$ diagrams (figure 10c–d), the majority of GPG samples fall in the field of reduced A-type granite, though some samples also fall in the oxidised A-type granite field. The samples of PG, on the contrary, fall in the field of oxidised A-type granites.

6.3 Petrogenesis

6.3.1 Implications from Trace and REEs

The Chondrite normalised REE patterns of GPG show less fractionated REE patterns, with enriched LREE and HREE and strong negative Eu anomaly (figure 8a). These features can be explained by constraining the source rock to a basic plagioclase rich-garnet free composition. Melting of such source

rock will result in melts with negative Eu anomalies and enriched HREE with low LREE/HREE ratios (Barker 1979; Martin 1999). These features can further be explained by the fractional crystallisation of plagioclase from the melt followed by late stage hornblende crystallisation to deplete the LREE and impart the low LREE/HREE ratios (Arth and Barker 1976; Barker 1979). PG, on the other hand, shows highly fractionated REE patterns (high LREE/HREE ratios) with weak negative to moderate positive Eu anomaly (figure 8c). The incompatible behaviour of Eu indicates melting of a source with little plagioclase, whereas the depletion of HREE is predominantly caused by low degrees of melting of garnet-rich sources which retain and deplete the melt in HREE (Drummond and Defant 1990; Martin *et al.* 2005). The other probable explanation for the depleted HREE in PG could be the late stage fractionation of heavy mineral phases like zircon, allanite, sphene, apatite and monazite because of their extremely high partition coefficients for HREE (Peter Gromet and

Figure 9. (a–f) Granitoid discrimination diagrams from Whalen *et al.* (1987) for GPG and PG samples. FG-field for fractionated I-type granitoids; OGT-field for I-, S- and M-type granitoids, (g–h) Y+Nb vs. Rb and Y vs. Nb tectonic discrimination diagrams of Pearce *et al.* (1984). Abbreviations of fields; VAG: volcanic arc granites; WPG: within plate granites; Syn-COLG: Syn-collisional granites; ORG: ocean ridge granites; Post-COLG: post-collision granites, (i) Zr concentrations in granites plotted against the cationic ratio M. The lines marked 700°–850°C indicate the proportions of Zr that can be dissolved in granitic melts of different compositions, expressed by the M parameter (Miller *et al.* 2003; Watson and Harrison 1983).

Silver 1983; Rollinson 1993). The Eu/Eu^* vs. SiO_2 plot (figure 7i) shows that the size of negative Eu anomalies decreases with increasing differentiation for GPG and *vice-versa* for PG. The reduction in size of the negative Eu anomalies for PG and concave HREE pattern, could possibly be attributed to hornblende fractionation because Eu has a lower partition coefficient than most MREE and HREE for hornblende in felsic rocks (Green 1994). High concentrations of Th, U and Pb in GPG and PG are indicative of the involvement of crustal components in their genesis (figure 8b and d). Negative anomalies of Nb, P and Ti are considered to be a characteristic feature of a subduction-related environment. The high values of Y for GPG, also suggest their formation by melting of a garnet

free source. Thus, the low Sr, high Y and HFSE contents with negative Eu anomaly indicate an origin at a shallower depth for GPG rocks (low P–T conditions) when compared to PG rocks which originate relatively at deeper levels.

6.3.2 Fractional crystallisation

The linear geochemical trends of major and trace elements illustrated by decreasing Al_2O_3 , CaO , MgO , $Fe_2O_3^t$, TiO_2 , P_2O_5 , Ba, Sr and Zr and increasing Rb and Rb/Sr with increasing silica coupled with the similarity and parallel nature of the normalised REE patterns and depletion in Ba, Sr, Nb, P, Ti and Eu in the spidergrams, suggest a

Figure 10. Whole-rock (a) $\text{CaO}/(\text{FeO}^t + \text{MgO} + \text{TiO}_2)$ vs. $\text{CaO} + \text{Al}_2\text{O}_3$, (b) $\text{CaO}/(\text{FeO}^t + \text{MgO} + \text{TiO}_2)$ vs. Al_2O_3 , (c) $\text{FeO}^t/(\text{FeO}^t + \text{MgO})$ vs. Al_2O_3 , and (d) $\text{FeO}^t/(\text{FeO}^t + \text{MgO})$ vs. $\text{Al}_2\text{O}_3/(\text{K}_2\text{O} + \text{Na}_2\text{O})$ discrimination diagrams with compositional fields from Dall'Agnol and de Oliveira (2007).

major role of crystal fractionation during the evolution of these rocks. The decrease in P_2O_5 , Fe_2O_3^t and TiO_2 with increasing SiO_2 (figure 6), along with strong negative Nb, P and Ti anomalies (figure 8b and d) suggests fractionation of apatite and Fe–Ti oxides for all samples. The depletion of U relative to Th is observed in the spidergrams of GPG and PG, indicating the fractionation of either zircon or allanite. The presence of negative Eu, Sr and Ba anomalies in these rocks are indicative of fractionation of both plagioclase and K-feldspar. The pronounced negative Eu anomaly in the REE patterns of GPG require extensive fractionation of plagioclase when compared to PG and is also supported by negative correlation of Al_2O_3 and CaO with SiO_2 (figure 6), and the positive one between SiO_2 vs. Rb/Sr and CaO vs. Sr (figure 7e and f). However, the increase of Ba with Sr (figure 7c) suggests a systematic removal of K-feldspar, biotite

and plagioclase from the melt for all samples. In addition, the fractionation of mafic phases particularly hornblende is suggested by the negative correlation of MgO and Fe_2O_3^t with SiO_2 (figure 6) and the positive correlation between MgO vs. Y (figure 7h) and A/CNK vs. SiO_2 (figure not shown) which is predominant in PG. The decreasing Zr content with increasing differentiation (figure 7g) for all samples in accordance with their zircon saturation temperatures indicates that with progressive fractionation, the solubility of Zr decreases in the melt because of falling temperature and, consequently, zircon is being removed from the melt (King *et al.* 1997).

6.3.3 Magma source

A number of petrogenetic schemes have been proposed for the origin of A-type granites involving

the crustal and mantle sources that reflect their distinct and diversified compositions (Kemp and Hawkesworth 2003). However, Loiselle and Wones (1979) originally suggested that A-type granites are formed by fractionation of mantle-derived alkali basalt with or without crustal interaction, whereas Eby (1990) showed that A-type granitic melts can also be formed from mixing between crustal and mantle sources. Therefore, three main modes of origin have been proposed by Frost and Frost (2011) on the basis of earlier studies to produce ferroan granitic compositions: (1) partial melting of quartz-feldspathic crustal rocks, (2) differentiation of tholeiitic or alkali basaltic magma, (3) a combination of the first two models, in which differentiating basaltic magmas assimilate crustal rocks.

From the above discussions, it is clear that GPG are ferroan, calc-alkalic, metaluminous to peraluminous reduced A-type granites though some samples show oxidised nature too. These features are consistent with their derivation from the low-pressure partial melting of quartz-feldspathic igneous sources with a metasedimentary rock contribution (Dall'Agnol and de Oliveira 2007), or alternatively from differentiation of tholeiitic sources with significant crustal contamination (Frost and Frost 2011). In contrast to GPG, less evolved PG granites are ferroan, alkali-calcic to calc-alkalic peraluminous showing oxidised A-type characteristics, which can be attributed to partial melting of lower crustal quartz-feldspathic igneous sources under relatively oxidising conditions with appreciable water contents (≥ 4 wt.%) (Dall'Agnol and de Oliveira 2007), or can be interpreted to form by partial melting of felsic granulite (Frost and Frost 2011). Eby (1990, 1992) has shown that the Y/Nb ratios are relatively constant and have very little or negligible effect during fractionation of A-type magmas. Therefore, this ratio can be regarded as a key to identify the sources of magma. Magmas with $Y/Nb < 1.2$ are derived from the mantle sources, whereas higher $Y/Nb (> 1.2)$ ratios imply solely crustal sources or mixed sources. The average Y/Nb ratios of GPG (2.70) and PG (1.52) are greater than 1.2, implying their derivation either from a crustal or mixed source.

The high-temperature nature of GPG and PG (T_{gr} : $\text{GPG}_{\text{avg}} \sim 861^\circ\text{C}$ and $\text{PG}_{\text{avg}} \sim 835^\circ\text{C}$) also suggest that they have been generated either in the lower crustal or upper mantle region. The heat required to melt the parent rocks at such high temperatures must be provided by the mantle-derived melts either directly or indirectly, i.e., they may only provide the necessary heat for crustal melting or they may contribute mass through differentiation and assimilation of crustal melts (Frost and Frost 2011). Also, the presence of the differentiated

basalt-andesite-rhyolite suite reported from the northern and eastern vicinity of Gaya (Ahmad and Wanjari 2009; Saikia *et al.* 2014; Wanjari *et al.* 2012) suggests an interaction of hot mafic magma with the crust.

To put further constraints on the three potential sources for A-type magmatism, we use the relationship of Y/Nb with Th/Nb, Th/Ta and Ce/Pb, as given by Moreno *et al.* (2014). These ratios provide an efficient discrimination between the continental crust, oceanic island and convergent margin rocks. On $(Y/Nb)_N$ vs. $(Th/Nb)_N$ diagram (figure 11a), all samples lie within the field of convergent margin magmatism plotting close to, or inside, the field of continental crust estimates. These relationships suggest an involvement of a continental crust component in the genesis of both GPG and PG which are formed in the convergent

Figure 11. (a) $(Y/Nb)_N$ vs. $(Th/Nb)_N$ and (b) $(Y/Nb)_N$ vs. $(Ce/Pb)_N$ discrimination diagrams with compositional fields after Moreno *et al.* (2014). Values normalised to the silicate earth after McDonough and Sun (1995).

margin, i.e., a collisional tectonic setting, also supported by negative Nb and positive Pb anomalies in spidergrams (figure 8b and d). Whereas on $(Y/Nb)_N$ vs. $(Ce/Pb)_N$ (figure 11b), all samples lie within the field of convergent margin magmatism but plot below the continental crust field. The significantly lower $(Ce/Pb)_N$ values presented by the granites suggest that this ratio was modified during magma differentiation. Thus, in the absence of isotopic constraints, the most likely origin for these granites seems to be the partial melting of crustal sources with variable mantle involvement, which is further evolved by the process of fractional crystallisation.

6.3.4 Tectonic setting: Anorogenic or post-orogenic?

A-type granites were originally defined as anorogenic granites by Loiselle and Wones (1979), but their exclusively anorogenic geotectonic settings

have been questioned and later studies suggested that they can be formed in both anorogenic and post-orogenic settings (Whalen *et al.* 1987; Eby 1992). Eby (1992) attempted to relate A-type granites with tectonic settings and further subdivided A-type into A_1 - and A_2 -type granites. According to this classification based on Y/Nb ratios, these two types represent different sources and tectonic settings, where A_1 ($Y/Nb < 1.2$) emplaced in an anorogenic setting, viz., continental rifts or intraplate environments and A_2 ($Y/Nb > 1.2$) in a post-orogenic setting, viz., continent–continent collision or island–arc setting. The Y/Nb ratios of all the studied samples vary from 1.10–4.69. Therefore, the granites of the study area plot in the A_2 -type granite field on Rb/Nb vs. Y/Nb and Y–Ce–Nb diagrams (figure 12a and b) suggest a post-collisional tectonic setting.

7. Conclusions

The granites present in the study area comprises grey and pink granites (GPG) which are Hbl-Bt rich whereas porphyritic granites (PG) are mainly Bt rich. Geochemically both the GPG and PG are ferroan, alkali-calcic to calc-alkalic, metaluminous to peraluminous, reduced to oxidised A-type granites. Coherent geochemical trends and similarities in the trace element behaviour along with certain differences in the REE pattern of GPG and PG suggest a key role of fractional crystallisation during petrogenesis of these granites. Trace element characteristics, REE and the tectonic discrimination diagrams reveal that they were derived from a predominant crustal source with a variable degree of mantle input. PG were produced from partial melting of quartz-feldspathic igneous sources under relatively oxidising conditions with appreciable water contents, emplaced at deeper levels than GPG. In comparison, GPG were emplaced at shallower levels and were produced from low-pressure partial melting of quartz-feldspathic igneous sources with a metasedimentary rock contribution with relatively low water activity. An A_2 -type affinity suggests that these granites are post-collisional but not anorogenic. Further constraining of the magma sources and the spatio-temporal relationship of the granites can only be revealed by studying isotopic ratios.

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Figure 12. (a) Y/Nb vs. Rb/Nb and (b) Y–Nb–Ce subdiscriminant diagrams for A-type granites from Eby (1992). A_1 : field for anorogenic A-type granites; A_2 : field for post-collisional A-type granites.

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