Contributions to the petrography, geochemistry and geochronology (U-Pb and Sm-Nd) of the Paleoproterozoic effusive rocks from Iricoumé Group, Amazonian Craton, Brazil

Contribuições para a petrografia, geoquímica e geocronologia (U-Pb e Sm-Nd) das rochas efusivas Paleoproterozóicas do Grupo Iricoumé, craton Amazônico, Brasil

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RESUMO: O sul do escudo das Guianas no craton Amazônico hospeda volumoso registro de rochas efusivas paleoproterozóicas do Grupo Iricoumé em notável estado de preservação de texturas e estruturas. O conteúdo em SiO₂ nessas rochas revela associação bimodal com rochas ácidas (SiO₂ > 67%) e intermediárias (SiO₂ < 57.7%). As rochas ácidas são riolitos a riodacitos com alto álcalis, Rb, Zr, Nb + Ta, La + Ce e 10⁴ Ga/Al, baixos Fe₂O_{3tot}, TiO₂, CaO, Sr e Co, exibindo composição subalcalina, metaluminosa a peraluminosa, geoquímica compatível ao magmatismo do tipo A pós-colisional a intra-placa. As rochas intermediárias são basalto andesíticos a andesitos com alto TiO₂, Al₂O₃, Fe₂O_{3total}, MgO, CaO, Sr e Co, baixo SiO₂, K₂O, Rb, Zr, Nb + Ta, La + Ce, composição subalcalina, metaluminosa e geoquímica de basalto intra-placa. A idade U-Pb de cristalização das rochas ácidas, obtida em zircão por LA-MC-ICPMS é de 1.882 ± 11 Ma. Os dados isotópicos Sm-Nd obtidos para os diferentes tipos de rochas revelaram idade modelo Nd T_{DM} entre 2.59 e 2.16 Ga e valores de ε_{Nd}(t) entre -5.78 e 0.03, indicando evolução magmática envolvendo retrabalhamento de crosta continental paleoproterozóica (crosta transamazônica) no período Rhyaciano-Sideriano, com algum grau de mistura com magmas derivados do manto ou contaminação por crosta arqueana. As informações petrográficas, geoquímicas e geocronológicas neste trabalho sugerem um ambiente pós-colisional a intra-placa para o vulcanismo Iricoumé, envolvendo elevação da crosta inferior e geração de magmas basálticos em um regime extensional.

PALAVRAS-CHAVE: cráton Amazônico; escudo das Guianas; Grupo Iricoumé; vulcanismo bimodal; Paleoproterozóico.

ABSTRACT: The southernmost region of the Guyana shield, Amazonian craton, hosts large record of Paleoproterozoic effusive rocks of the Iricoumé Group. They present remarkably well-preserved igneous textures and structures. The SiO₂ contents reveal a bimodal association marked by a compositional gap between acid (SiO₂ > 67 wt%) and intermediate (SiO₂ < 57.7 wt%) rocks. The acid effusive rocks are rhyolites to rhyodacites with high SiO₂, alkali, Rb, Zr, Nb + Ta, La + Ce and 10⁴ Ga/Al content and low Fe₂O_{3wt}, TiO₂, CaO, Sr and Co content. They exhibit subalkaline, metaluminous-to-peraluminous compositions, and geochemically compatible to A-type magmatism emplaced in post-collisional to within-plate tectonic settings. The intermediate rocks are andesitic/basalt to andesite relatively high contents of TiO₂, Al₂O₃, Fe₂O_{3total}, MgO, CaO, Sr and Co; low SiO₂, K₂O, Rb, Zr, Nb + Ta, La + Ce. They have subalkaline and metaluminous geochemical composition and plot on within-plate basalt field. The acid rocks crystallized at 1882 ± 11 Ma in U-Pb analyses for LA-MC-ICPMS zircon data. The Sm-Nd isotopic data on all rocks reveal a Nd TDM model ages between 2.59 and 2.16 Ga and ε_{Nd}(t) values between -5.78 and 0.03, indicate that the magmatic evolution was related to the reworking of older Paleoproterozoic at the Rhyacian-Siderian period, continental crust (Transamazonian crust-forming event) with some mixing with a limited amount mantle-derived magmas or with contamination by Archean crust. The petrographic, geochemical and geochronological data presented in this paper suggest a within-plate to post-collisional tectonic setting for the Iricoumé volcanism, involving lower crust uplift and generation of basalt magma in an extensional regime.

KEYWORDS: Amazonian craton; Guyana shield; Iricoumé Group; bimodal volcanism; Paleoproterozoic.

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INTRODUCTION

The central part of the Amazonian craton hosts extensive records of Paleoproterozoic volcano-plutonic magmatic associations distributed on NW–SE trend (Fig. 1A). They are included in regional units known as Iricoumé–Mapuera within the Guyana shield and Iri-Maloquinha, within the Guaporé shield, comprising the voluminous Uatumá magmatic event (Tassinari & Macambira 1999; Tassinari *et al.* 2000; Santos *et al.* 2000; Pinho *et al.* 2004; CPRM 2006 & 2008; Fernandes *et al.* 2006; Ferron *et al.* 2006 & 2010; Valério *et al.* 2009 & 2012; Juliani & Fernandes 2010; Rocha *et al.* 2012; Klein *et al.* 2012).

The Iricoumé–Mapuera magmatic association consists of effusive and pyroclastic rock units of acid to intermediate composition forming the Iricoumé Group, which is intruded by geochemically similar alkali-rich granites known as the Mapuera Intrusive Suite. The northeastern region of Amazonas state (Fig. 1) displays excellent exposures of the Iricoumé–Mapuera association, which, during the last decade, have become the object of scientific research focusing on the petrographic, geochemical and geochronological characteristics (Valério 2006, 2011; Ferron 2006; Pierosan 2009; Marques 2011; Costa 2010). These studies have demonstrated that the Iricoumé–Mapuera association is formed by A-type magmatism with metaluminous to weakly peraluminous compositions, with geochemical affinity towards within-plate to post-collisional tectonic setting. The available geochronological data, mainly those obtained from the zircon $^{207}\text{Pb}/^{206}\text{Pb}$ evaporation method, indicate crystallization ages between 1879 and 1897 Ma. However, Nd–Sr–Hf isotopic data are still limited or nonexistent. Tab. 1 summarizes the isotopic information available for the Iricoumé volcanic rocks in the southern Guyana shield.

In this study, we present new data on the petrography, whole-rock geochemistry and geochronology (U–Pb and Sm–Nd) of effusive rocks from the Iricoumé Group in the northeastern region of Amazonas state, and compare them with the information available in the literature. Their petrogenesis and the geodynamic evolution associated with the Paleoproterozoic volcanic processes in the Amazonian craton.

ANALYTICAL METHODS

The petrographic investigations were performed at the microscopy laboratory of the Department of Geosciences, Universidade Federal do Amazonas (DEGEO-UFAM). The samples were chosen and prepared for geochemistry and isotopic (Sm–Nd) analyses at the laboratories of the DEGEO-UFAM, using conventional crushing in a jaw mill and in agate pot. Geochemical analyses were undertaken at

the ACME Analytical Laboratories, Vancouver, Canada. The samples were analyzed for major elements (SiO_2 , TiO_2 , Al_2O_3 , $\text{Fe}_2\text{O}_{3\text{tot}}$, MnO , MgO , CaO , Na_2O , K_2O and P_2O_5) by Inductively Coupled Plasma-Emission Spectrometry and for trace and rare-earth elements by Inductively Coupled Plasma-Mass Spectrometry (ICPMS).

The isotopic analyses (U–Pb and Sm–Nd) were carried out at the isotope geology laboratories of the Universidade de Brasília, by applying the analytical procedures described below:

- U–Pb analyses by LA-MC-ICPMS followed the analytical procedure described by Bühn *et al.* (2009). Zircon concentrates were extracted using conventional gravimetric and magnetic separation techniques. The zircon grains were selected under a binocular microscope to obtain fractions of similar size, shape and colour. For *in situ* U–Pb isotopic analysis, hand-picked zircon grains were mounted in epoxy mounts and polished to obtain a smooth surface. The samples were mounted in an in-house, specially designed laser cell and loaded into a New Wave UP213 Nd:YAG laser ($\lambda = 213 \text{ nm}$), linked to a Thermo Finnigan Neptune Multi-collector ICPMS. Helium was used as the carrier gas and mixed with argon before entering the ICP. The laser was run at a frequency of 10 Hz and energy of $\sim 100 \text{ mJ/cm}^2$ with a spot size of $30 \mu\text{m}$ for U–Pb dating and $40 \mu\text{m}$ for Hf isotopic analyses. Plotting of U–Pb data and age calculations were performed using ISOPLOT 3.0 (Ludwig 2003), and errors for isotopic ratios are presented at 1σ level.
- Sm–Nd isotopic analyses followed the method described by Gioia & Pimentel (2000). Whole-rock powders (ca. 50 mg) were mixed with a ^{149}Sm – ^{150}Nd spike solution and dissolved in Savillex capsules. Sm and Nd extraction of whole-rock samples followed conventional cation exchange techniques, using Teflon columns containing LN-Spec resin ([di-(2-ethylhexyl) phosphoric acid (HDEHP) supported on PTFE powder). Sm and Nd samples were loaded on re-evaporation filaments of double filament assemblies and the isotopic measurements were carried out on a multi-collector Finnigan MAT 262 mass spectrometer in static mode. Uncertainties for Sm/Nd and $^{143}\text{Nd}/^{144}\text{Nd}$ ratios are better than $\pm 0.2\%$ (2σ) and $\pm 0.003\%$ (2σ), respectively, based on repeated analyses of international rock standards BHVO-1 and BCR-1. $^{143}\text{Nd}/^{144}\text{Nd}$ ratios were normalized to $^{146}\text{Nd}/^{144}\text{Nd}$ of 0.7219 and the decay constant (λ) used was 6.54×10^{-12} .

GEOLOGICAL SETTING

The southernmost of the Guyana shield includes the boundary region between the Paleoproterozoic basement and

Figure 1. (A) Distribution of the Uatumã magmatic event in the Brazilian portion of the Amazonian craton and location of the study area (modified from Pierosan *et al.* 2011a). (B) Regional geological map and localization of the study area (modified from Valério *et al.* 2012). (C) Geological map of the study area (modified from Marques 2011).

the Silurian-Devonian to Cretaceous sedimentary cover of the northern edge of the Amazon Basin (Fig. 1). The Paleoproterozoic basement includes gneisses, migmatites, meta-granites, amphibolites and charnockites comprising the so-called Jauaperi/Anauá Complex. The geochronological data on the basement rocks are still inconclusive and vary from 2.02 to 1.86 Ga (Santos *et al.* 2002; Reis *et al.* 2003; CPRM 2006; Almeida 2006). In the study area, the nature of these metamorphic rocks is controversial. Some geologists believe that they represent products of contact metamorphism produced by Paleoproterozoic Mapuera Intrusive Suite, represented by the A-type São Gabriel batholith, but others suggest that they are simply the exposure of Paleoproterozoic basement rocks (Fig. 1C).

Four Paleoproterozoic (1900 – 1782 Ma) involving I- and A-type magmatic events are recognized in the region (Fig. 1B) as follows: (1) the Água Branca Intrusive Suite (1900 – 1890 Ma) is characterized by an expanded granite series that varies from predominantly porphyritic to seriate-textured monzogranites and granodiorites to diorites. These rocks are metaluminous to peraluminous, calc-alkaline with high-K I-type geochemical signature (Araújo Neto & Moreira 1976; Almeida 2006; CPRM 2006; Almeida & Macambira 2007; Valério 2006, 2011; Valério *et al.* 2006, 2009, 2012); (2) the Iricoumé Group (1897 – 1881 Ma) comprises effusive rocks and pyroclastic flow associations. These rocks are metaluminous to weakly peraluminous, subalkaline with high-K and A-type affinity; (3) the Mapuera Intrusive Suite (1889 – 1871 Ma) is represented by monzogranite, syenogranite and alkali-feldspar granite stocks. These rocks are weakly peraluminous, subalkaline with high-K, and have geochemical characteristics similar to A-type granites. Due to the geochemical and geochronological similarities between felsic volcanic rocks of the Iricoumé Group and granitic rocks of the Mapuera Suite, these rocks have been interpreted as a coeval volcano-plutonic association and inserted as part of the regional units known as the Uatumã Supergroup (Oliveira *et al.* 1975; Araújo

Neto & Moreira 1976; Veiga *et al.* 1979; Schobbenhaus *et al.* 1994; Costi *et al.* 2000; CPRM 2006; Valério *et al.* 2009, 2012; Ferron *et al.* 2006, 2010; Pierosan *et al.* 2011a, 2011b; Berreto *et al.* 2013); and (4) the Madeira Intrusive Suite (1834 – 1782 Ma) occurs in the Pitinga Sn-mine area and is composed by biotite-syenogranite, alkali-feldspar rapakivi, hipersolvus and albite granite stocks bearing cryolite-zircon-cassiterite-pyroxene-columbite-xenotime mineralization. These rocks are peraluminous to metaluminous, rarely peralkaline, and also present geochemical and mineralogical characteristics of A-type granites (Horbe *et al.* 1991; Fuck *et al.* 1993; Lenharo 1998; Costi *et al.* 2000, 2005; Ferron *et al.* 2006, 2010). The Paleoproterozoic geological record is completed by the volcano-sedimentary cover of the Urupi Formation and by the ca. 1.79 Ga old mafic-tholeiitic rocks of the Quarenta Ilhas Formation (CPRM 2006). In addition, a magmatic event marked by gabbroic intrusions is also recognized in the region but has not been studied in detail yet (Souza & Nogueira 2009).

The Meso- to Neoproterozoic geological context is represented by the alkaline magmatism of the Seringa Formation (1.07 Ga) and by siliciclastic sedimentary sequence of the Prosperança Formation (CPRM 2006; Cunha *et al.* 1994; Nogueira 1999), while the Phanerozoic Trombetas and Javari Groups represent the Amazon sedimentary basin (Cunha *et al.* 1994). In addition, Mesozoic diabase dikes and Cenozoic laterite cover occur in the area as well (CPRM 2006; Horbe *et al.* 2001).

Effusive Rocks of the Iricoumé Group

The Iricoumé Group exposures are partially covered by dense rainforest, thick soil cover, by the lake of the Balbina hydroelectric dam, as well as by restricted access areas (e.g. indigenous reserves, Fig. 1B). However, in the farms and agricultural settlement areas, as well as in the mining fronts, excellent exposures of the Iricoumé volcanism are observed, with remarkable preservation of original igneous textures and structures.

Table 1. Isotopic resume available for the Iricoumé volcanism in the southern Guyana shield

Region	Iricoumé volcanism lithologies	Age (Ma)	Method	$\epsilon_{\text{Nd}}(t)$	$\text{NdT}_{\text{DM}}(\text{Ga})$	References
SE Roraima state	dacite	1893	zircon $^{207}\text{Pb}/^{206}\text{Pb}$ by evaporation	-	-	Macambira <i>et al.</i> (2002)
Pitinga mining distric (Amazonas state)	rhyolite, rhyodacites, ignimbrite	1890 – 1881	zircon $^{207}\text{Pb}/^{206}\text{Pb}$ by evaporation	-	-	Costi <i>et al.</i> (2000), Costa (2010), Ferron (2006)
NE Amazonas state	rhyolite, dacite, rhyodacites, ignimbrite	1896 – 1883	zircon $^{207}\text{Pb}/^{206}\text{Pb}$ by evaporation	1.69 to -1.44	2.26 to 2.05	CRPM (2006), Valério (2006 and 2011)
NE Amazonas state	rhyolite, rhyodacites, andesite, tracky-andesite	1882	zircon U-Pb by LA-MC-ICPMS	- 5.43 to 0.80	2.59 to 2.16	This paper
NW Pará state	andesite, ignimbrite	1992 – 1888	zircon $^{207}\text{Pb}/^{206}\text{Pb}$ by evaporation	-3.0 to -2.3	1.98 to 2.39	Barreto <i>et al.</i> (2012 and 2013) Klein <i>et al.</i> (2012)

Petrography

The acid rocks are rhyolites to rhyodacites, which exhibit red to pink-gray colour, fine to medium-grained porphyritic texture marked by feldspars and quartz-phenocrysts embedded in a fine-grained matrix. Amygdales and fractures filled by late- to post-magmatic minerals, such as epidote, chlorite, carbonate and quartz, are also recognized. Occasionally, the outcrops display magmatic layering with sense of flow to SW (Fig. 2A). The absence of vitroclastic and lithoclastic textures, associated with myrmekite features locally present in these rocks, indicates that these are sub-aerial to aerial lava-flows. However, in some outcrops displaying subhorizontal to curvy-planar layering, where occasional pumice, lithic and fiamme are observed, suggesting that these represent ignimbrite-flows modified by high-temperature rheomorphic process (e.g. Pioli & Rosi 2005; Andrews & Branney 2011), which makes it difficult to distinguish between ignimbrite-flows and lava-flows.

In general, the acid rocks present modal contents of alkali-feldspar, plagioclase (oligoclase, An_{26-30}) and quartz-phenocrysts varying from 15 to 60 vol.%. They display euhedral to subhedral shapes, with lengths ranging from 0.3 to 3 mm, and commonly exhibit corrosion and resorption features, undulating extinction and are low to moderately fractured (Fig. 2B and C). Bulbous-type myrmekite is a common occurrence in the rims of larger crystals. Quartz-feldspar groundmass presents fine-grained phaneritic to aphanitic textures, with crystals usually oriented in the magmatic-flow circumventing the phenocrysts. They also have subhedral crystals of hornblende, biotite, titanite, zircon, hematite, magnetite, pyrite, ilmenite, allanite and apatite as accessory. Hornblende and biotite crystals are partially chloritized as phenocrysts shapes with lengths ranging from 0.3 to 0.8 mm in the finer groundmass. Frequently, ilmenite crystals involved by a thin rim of rutile are observed, and may probably represent the product of Fe-Ti excess exsolved from the ilmenite structure during temperature decrease or oxygen fugacity variation (e.g. Haggerty 1991).

The intermediate rocks are andesite to trachy-andesite, which exhibit green to green-greyish colour, fine-grained porphyritic texture highlighted by plagioclase-phenocrysts embedded in a fine-grained to aphanitic matrix. The outcrops also display subhorizontal layering marking the magmatic flow and frequent amygdales filled by late- to post-magmatic minerals, such as epidote, chlorite, carbonate and quartz (Fig. 2D). The phenocrysts, ranging from 25 to 50 vol.%, are mainly plagioclase (andesine, An_{38-46}), and rare alkali-feldspar. Plagioclase-phenocrysts have euhedral to subhedral prismatic shapes, with lengths ranging from 1 to 15 mm usually forming a glomeroporphyritic texture (Fig. 2E). They exhibit albite-carlsbad twinning, are slightly

fractured and have corroded edges, moderate decalcification stage marked by epidote and white-mica generation. The groundmass has a fine phaneritic texture and is composed of short lath-shaped plagioclase crystals with subophitic to trachytic texture oriented along the magmatic-flow, circumventing the phenocrysts. They also have partially chloritized hornblende and biotite, as well as pyroxene crystals partially replaced by fibrous amphibole (actinolite-tremolite) form interstitial mineral arrangement. Quartz, titanite, magnetite, ilmenite, pyrite, chalcopyrite and apatite are present as accessory (Fig. 2F).

Whole-Rock Geochemistry

Whole-rock compositions for acid to intermediate rocks of the Iricoumé Group are shown in Tab. 2 and are discussed below.

The acid rocks present the following contents: $SiO_2 = 67 - 72.19$ wt%, $Al_2O_3 = 12.94 - 14.81$ wt%, $Fe_2O_{3total} = 3.37 - 4.07$ wt%, $CaO = 0.6 - 2.59$ wt%, $Na_2O = 3.25 - 3.88$ wt%, $K_2O = 4 - 6$ wt%, low contents of TiO_2 , MnO, MgO, P_2O_5 and $LOI < 1$ wt%, with $Na_2O + K_2O$ between 7.75 and 9.39 wt%, K_2O/Na_2O ratio ≥ 1 , and $FeO_t/FeO_c + MgO$ ratio ranging from 0.78 to 0.95. They present high Ba, Rb, Zr, HREE, and low Sr, Ta, Co and Hf contents. On the $Na_2O + K_2O$ versus SiO_2 (TAS), as well as Zr/TiO_2 versus SiO_2 and Zr/TiO_2 versus Nb/Y diagrams, the analyses plot on the rhyolite to rhyodacite fields (Fig. 3), with high alkali contents indicating high-K subalkaline compositions. On the A/NK versus A/CNK diagram, this sample plots between metaluminous to peraluminous fields (Fig. 3D).

On the multi-element spidergrams, these rocks display a pattern similar to upper continental crust, marked by positive Rb, Th, K, La, Ce, Nd and Zr anomalies and strong negative Ba, Nb, Sr, P and Ti anomalies (Fig. 3E; e.g. Thompson 1982; Rollinson 1993). The REE-normalized patterns are characterized by high LREE ($La + Ce = 157 - 226$) and relatively depleted HREE ($Tm + Yb = 4.82 - 6.76$) contents in relation to the chondrite (Fig. 3G). The REE distribution patterns present moderate fractionation of the LREE group with $(La/Sm)_N = 4.11 - 5.26$ and absence or light-fractionation of the HRRE group with $(Gd/Yb)_N = 1.24 - 1.32$, separated by negative Eu anomaly ($Eu/Eu^* = 0.21 - 0.28$).

The acid rocks of the Iricoumé Group have been recognized as A-type (Valério *et al.* 2009; Ferron *et al.* 2010; Pierosan *et al.* 2011a). In the study area, these rocks also present geochemical attributes compatible with A-type magmatism, such as relatively high alkalis ($Na_2O + K_2O$), Zr, Th, Nb and La + Ce contents, as well as high $10^4 Ga/Al = 2.35 - 2.90$ ratios (Collins *et al.* 1982; Clemens *et al.* 1986; Whalen *et al.* 1987; Eby 1992). On the Zr and $Zr + Nb + Ce + Y$ versus $10^4 Ga/Al$ diagrams, the samples plot in the

A-type granite field (Fig. 4A and 4B), and on the Nb–Y–Ce diagram, the samples plot in the A_2 -subtype field (Fig. 4C), with Y/Nb ratios > 1.2, indicative of magmas emplaced during extensional collapse according to Eby (1992). These

geochemical characteristics are consistent with the information obtained about the Iricoumé volcanism in the Pitinga mining district, located about 150 km to the north of the study area (Fig. 1B).

Figure 2. Macroscopic and microscopic features of the effusive rocks of the Iricoumé Group. (A) Magmatic flow layering with sense of flow to SW in acid rocks; (B and C) porphyritic texture in thin section on rhyolites and rhyodacites, respectively; (D) amygdaloid and fractures filled by late- to post-magmatic minerals in andesites; (E and F) porphyritic to glomeroporphyritic micro-texture with amygdaloid in andesites.

The intermediate rocks present the following contents: $\text{SiO}_2 = 55.91 - 57.69$ wt%, $\text{Al}_2\text{O}_3 = 16.66 - 15.85$ wt%, $\text{Fe}_2\text{O}_{3\text{total}} = 7 - 10.64$ wt%, $\text{MgO} = 1.13 - 4.47$ wt%, $\text{CaO} = 5.68 - 7.21$ wt%, $\text{Na}_2\text{O} = 2.69 - 3.34$ wt%, $\text{K}_2\text{O} = 2.23 - 3.27$ wt%, low contents of TiO_2 and $\text{P}_2\text{O}_5 < 1.3$ wt%, and $\text{LOI} = 0.80 - 2.40$ wt%, with $\text{Na}_2\text{O} + \text{K}_2\text{O}$ between 5.47 and 6.54 wt%, $\text{K}_2\text{O}/\text{Na}_2\text{O}$ ratios are ≤ 1 , and $\text{FeO}_t/\text{FeO}_t +$

MgO values range from 0.61 to 0.89. They display relatively high Ba, Sr, Zr, Co and HREE contents and low Rb, Ta, Th and Hf contents. On the $\text{Na}_2\text{O} + \text{K}_2\text{O}$ versus SiO_2 (TAS) diagram the samples plot at the interface between the trachy-andesite, andesite/basaltic and andesite fields, indicating a bimodal magmatism, whose alkali content defines subalkaline composition (Fig. 3A). It is very likely

Table 2. Whole-rock geochemistry composition of the effusive rocks of the Iricoumé Group

Lithologies	Andesite to tracky-andesite									Rhyolite to rhyodacite						
Samples	SVP3A	SVP3A2	SVP4A	SVP36	SVP2E	SVP2E2	SVP2F	SVP3B	C6204	SVP20	SVP3C1	GC3753	C6604	GC3735	GC3717	GC3715
SiO_2 (wt%)	55.91	56.12	56.20	56.37	56.58	56.76	57	57.32	57.69	67.04	70.36	70.57	70.57	71	71.12	72.19
TiO_2	1.20	1.27	0.67	1.14	0.71	0.92	0.68	1.21	1.20	0.60	0.43	0.26	0.39	0.26	0.26	0.24
Al_2O_3	16.35	16.34	16.44	16.66	16.31	15.85	16.52	16.42	16.16	14.81	13.63	13.63	14.16	13.35	13.24	12.94
$\text{Fe}_2\text{O}_{3\text{total}}$	10.64	10.42	7.30	10.43	7.36	9.23	7	9.73	10.43	4.07	3.58	4.04	3.37	4.05	4.01	3.56
MnO	0.22	0.16	0.11	0.19	0.10	0.13	0.10	0.15	0.18	0.08	0.07	0.11	0.07	0.07	0.10	0.09
MgO	1.32	1.29	4.47	2.07	4.47	2.99	4.23	1.86	1.13	0.97	0.89	0.18	0.88	0.18	0.29	0.14
CaO	7.01	7.21	6.27	5.88	6.79	5.68	6.71	6.22	6.35	2.59	2.14	1.23	1.45	0.79	0.81	0.60
Na_2O	2.69	2.76	2.98	2.96	2.80	3.27	2.88	3.01	3.34	3.88	3.83	3.66	3.67	3.81	3.25	3.81
K_2O	2.78	2.84	2.65	2.92	2.62	3.27	2.74	2.92	2.23	4.31	4.01	5.57	4.08	5.58	6.04	5.21
P_2O_5	0.19	0.19	0.16	0.17	0.18	0.20	0.17	0.19	0.20	0.15	0.14	0.03	0.12	0.03	0.03	0.04
LOI	1.30	1.20	2.40	0.80	2	1.50	1.90	0.80	0.80	1.10	0.80	0.50	0.50	0.50	0.40	0.80
Total	99.61	99.80	99.65	99.59	99.93	99.80	99.83	99.83	99.71	99.63	99.88	99.58	99.26	99.62	99.55	99.62
Ba (ppm)	986	914	718	893	703.30	833.30	701.10	718.80	965.40	1240	536.50	790	531.40	600	801	681
Rb	79.90	80.60	116.90	88.70	88.20	129	97.80	84.20	82	152.50	137.20	216.60	130.90	189.40	230	194.60
Sr	257.10	251.90	344.40	278.50	362.20	274.40	352.70	283.30	170.50	238.20	185.60	95.10	192.60	95.20	77.70	73.50
Y	42.90	48.80	26.10	40	27.90	40	28.20	45.50	42.10	33	42.80	54.80	40.30	58.60	55.30	49.30
Zr	271.70	266.10	221.20	265.60	207.50	257.50	214.50	251.90	248.20	335.70	368.50	476.70	329.80	447.70	472.80	430
Nb	12.60	11.50	11.90	11.90	10.90	12.40	10.80	10.80	12	16.40	17.20	21.80	15.20	22.20	22.60	21.40
Ta	0.80	0.80	0.80	0.70	0.80	1	1.10	0.80	0.70	1.30	1.40	1.40	1.20	1.50	1.50	1.30
Co	46.90	54.70	43.20	30.70	37.10	44.20	41.50	46.80	25.60	127.30	46	3.40	5.70	4.10	3.80	4.60
Ga	21.40	21.30	16.60	20.80	16.81	18.50	18.10	21.10	22.10	16.20	19.60	18.80	20	17.80	18.70	18.10
Hf	7	7.50	5.20	6.90	6.30	7.50	5.50	7.40	6.90	9.20	9.40	13.40	9.40	12.60	13.10	11.80
Th	8.50	10.30	10.10	8.50	12.30	12.60	11.70	7.60	9.90	17.10	19.80	23.70	18.80	25.40	25.50	22.40
U	2.80	2.70	3.80	2.60	3.50	3.30	3.70	2.60	2.10	5.60	5.10	7.30	4.90	6.40	9.20	3.10
Cu	29.70	41.40	94.90	8.50	125.50	140.80	126.60	7.10	5	5.50	11.40	7.20	5.80	4.20	6.10	20.60
Zn	69	80	47	58	49	90	55	46	90	51	92	77	75	27	63	66
La	40.30	44.40	32.70	37.50	34	45	34	40.20	38.50	49.30	74.50	69.30	69	71.70	70.30	65.40
Ce	89.60	93	73	88.60	74.50	94.90	74.50	88.60	82.30	108.10	150.70	150.10	136.90	154.40	150.20	138.20
Pr	10.10	10.89	7.81	9.63	8.17	10.25	8.17	10.20	9.47	11.17	15.93	16.80	15.16	17.31	16.77	15.43
Nd	41.40	42.60	31.20	41.20	29.40	37.80	29.40	38.90	39.10	42.10	55.40	63.10	56.30	65.80	62.40	58.40
Sm	7.72	8.10	4.92	7.50	5.50	7	5.50	7.70	7.90	6.74	8.90	10.34	8.90	10.94	10.41	9.46
Eu	1.85	1.98	1.01	1.90	0.97	1.49	0.97	1.83	1.86	1.27	0.95	1.34	1.05	1.29	1.25	1.20
Gd	7.40	7.38	4.53	7.33	4.25	6.14	4.25	7.37	7.73	5.79	6.39	9.31	6.96	9.33	9.11	8.15
Tb	1.30	1.34	0.74	1.28	0.93	1.05	0.93	1.35	1.34	0.96	1.31	1.57	1.21	1.59	1.57	1.43
Dy	7.49	8.10	4.23	7.38	2.26	6.65	4.26	7.27	7.49	5.69	6.65	8.63	6.87	8.85	8.77	8.06
Ho	1.55	1.67	0.89	1.47	0.90	1.31	0.90	1.52	1.47	1.11	1.37	1.80	1.41	1.88	1.86	1.67
Er	4.39	4.68	2.73	4.48	2.71	4.16	2.71	4.09	4.38	3.49	4.30	5.53	3.95	5.79	5.73	5.26
Tm	0.69	0.77	0.40	0.64	0.45	0.58	0.45	0.69	0.64	0.56	0.66	0.89	0.65	0.92	0.88	0.78
Yb	4.31	4.17	2.68	4.33	2.39	3.44	2.39	4.23	4.26	3.77	4.18	5.68	4.17	5.84	5.75	5.17
Lu	0.63	0.73	0.39	0.63	0.45	0.54	0.45	0.65	0.63	0.57	0.67	0.88	0.68	0.93	0.88	0.79
$\text{Na}_2\text{O} + \text{K}_2\text{O}$	5.47	5.60	5.63	5.88	5.42	6.54	5.62	5.93	5.57	8.19	7.84	9.23	7.75	9.39	9.29	9.02
$\text{K}_2\text{O}/\text{Na}_2\text{O}$	1.03	1.02	0.89	0.98	0.93	1	0.95	0.97	0.66	1.11	1.04	1.52	1.11	1.46	1.85	1.36
$\text{FeO}_t/\text{FeO}_t + \text{MgO}$	0.87	0.87	0.61	0.82	0.61	0.74	0.61	0.82	0.89	0.79	0.79	0.94	0.78	0.94	0.92	0.95
$(\text{La}/\text{Sm})_N$	3.29	3.44	4.18	3.14	3.88	4.04	3.88	3.28	3.06	4.60	5.26	4.21	4.87	4.11	4.24	4.34
$(\text{Gd}/\text{Yb})_N$	1.39	1.43	1.37	1.37	1.44	1.44	1.44	1.41	1.47	1.24	1.23	1.32	1.35	1.29	1.28	1.27
Eu/Eu^*	0.41	0.44	0.22	0.42	0.21	0.33	0.21	0.40	0.41	0.28	0.21	0.29	0.23	0.28	0.27	0.26
$10^4\text{Ga}/\text{Al}$	3.10	3.09	2.41	3.02	2.44	2.68	2.62	3.06	3.20	2.35	2.84	2.73	2.90	2.58	2.71	2.62
$\text{Zr} + \text{Nb} + \text{Ce} + \text{Y}$	416.80	420.50	332.20	406.10	320.80	107.30	328	396.80	384.60	493.30	579.20	703.4	522.20	682.90	700.90	638.90

Figure 3. Geochemical characteristics of the volcanic rocks of the Iricoumé Group. (A) $\text{Na}_2\text{O} + \text{K}_2\text{O}$ versus SiO_2 diagram (Le Maitre *et al.* 1989); (B and C) Zr/TiO_2 versus Nb/Y and Zr/TiO_2 versus SiO_2 diagrams (Winchester & Floyd 1977); (D) alumina saturation index diagram (Shand indexes in Maniar & Piccoli 1989); (E and F) multi-element spidergrams for acid and intermediate rocks (chondrite-normalized pattern of Thompson 1982); (G and H) REE-normalized patterns diagrams for acid and intermediate rocks (chondrite-normalized pattern of Boynton 1984).

that the chemical compositions of major-elements have been modified by the presence of amygdaloidal texture. In this case, minor- and trace-element correlation diagrams are suitable. Thus, on the Zr/TiO_2 versus SiO_2 and Zr/TiO_2

versus Nb/Y diagrams, these rocks tend to plot in andesite to trachy-andesite fields (Fig. 3B and 3C). On the A/NK versus A/CNK diagram, this sample plots in the metaluminous field (Fig. 3D).

In the multi-element spidergrams, these rocks also demonstrate a pattern similar to upper continental crust, marked by positive Rb, Th, K, La, Ce, Nd, Sm, Zr and Hf anomalies and negative Ba, Nb, Ta, Sr, P and Ti anomalies (Fig. 3F, e.g. Thompson 1982; Rollinson 1993). The REE-normalized patterns are characterized by moderate LREE ($\text{La} + \text{Ce} = 105 - 140$) and relatively depleted HREE ($\text{Tm} + \text{Yb} = 2.80 - 5$) contents in relation to the chondrite (Fig. 3H). The REE distribution patterns also present moderate fractionation of the LREE group with $(\text{La}/\text{Sm})_{\text{N}} = 3.06 - 4.18$ and absence or light-fractionation of the HREE group with $(\text{Gd}/\text{Yb})_{\text{N}} = 1.37 - 1.47$, separated by negative Eu anomaly ($\text{Eu}/\text{Eu}^* = 0.21 - 0.44$).

Geochronology (U-Pb and Sm-Nd)

U-Pb and Sm-Nd data for the volcanic rocks of the Iricoumé Group are listed in Tabs. 3 and 4, and are discussed below.

The rhyolite SVP3C sample (810915/9823114 UTM coordinates) was chosen for U-Pb by LA-MC-ICPMS isotopic analyses and a total of 28 zircon crystals were handpicked. Zircon crystals are euhedral to subhedral, pale yellow to pale pink, zoned and displaying short- to long-prismatic shape (180 – 500 μm), with few to moderate micro-inclusions or fractures. Typical fine oscillatory zoning may be observed (Fig. 5). The U-Pb results obtained for 21 analyzed zircon crystals allowed the calculation of a discordia line indicating an upper intercept age of 1882 ± 11 Ma, which is here interpreted as the crystallization age of the rhyolite (Fig. 5). These data are consistent with the Pb–Pb interval age (1881 – 1896 Ma) known for the Iricoumé volcanism in this region (Tab. 1; e.g. Macambira *et al.* 2002; Valério 2006; Ferron 2006; Berreto *et al.* 2013).

The whole-rock Sm-Nd analyses were performed on seven selected samples, three acid (SVP3C1, GC3735, C6604) and four intermediate volcanic rocks samples (SVP2E, SVP4A, SVP3A, SVP3B).

As all samples showed $^{147}\text{Sm}/^{144}\text{Nd}$ ratios in the range of 0.10 – 0.12, these data were useful for calculation of T_{DM} model ages. For the acid rock samples, the $\epsilon_{\text{Nd}}(t = 1882 \text{ Ma})$ values ranged from -4.43 to 0.03, while $\text{Nd } T_{\text{DM}}$ model ages ranged from 2.59 to 2.16 Ga, and fractionation indexes ($f_{\text{Sm}/\text{Nd}}$) between -0.38 and -0.51. The $f_{\text{Sm}/\text{Nd}}$ values were calculated using the Goldstein *et al.* (1984) equation. For the intermediate rock samples, for which zircon ages are not available, the age of the rhyolite was assumed as their crystallization age, based on stratigraphic correlations. The $\epsilon_{\text{Nd}}(t = 1882 \text{ Ma})$ between -3.74 and 0.80, whereas T_{DM}

Figure 4. Granitoid discriminating diagram applied to acid rocks of the Iricoumé Group. (A and B) Zr and $\text{Zr} + \text{Nb} + \text{Ce} + \text{Y}$ versus $10^4 \text{Ga}/\text{Al}$ plot (Whalen *et al.* 1987); (C) Nb versus Y versus Ce triangular diagram for A-type granitoids (Eby 1992).

model ages are between 2.44 and 2.13 Ga. The fractionation indexes ($f_{\text{Sm/Nd}}$) are between -0.34 and -0.45. This data is also consistent with the Sm-Nd data known for Iricoumé volcanism in this region (Tab. 1).

The $\text{Nd } T_{\text{DM}}$ model age values for all samples are between 2.59 and 2.16 Ga. The $\epsilon_{\text{Nd}}(t)$ values from -4.78 to 0.03 indicate that the magmatic evolution was dominantly produced by related ensialic reworking of older Paleoproterozoic crust, with some degree of mixing of the felsic magmas with Paleoproterozoic (1.88 Ga) mantle-derived magmas. The diagram ϵ_{Nd} versus time summarizes the Nd evolution patterns provided by the rocks samples investigated (Fig. 6), and it suggests the existence of two isotopically distinct groups: (1) the group (C6604, SVP2E, SVP3A, SVP3B) display $\epsilon_{\text{Nd}}(t)$ values close to zero, suggesting a more contribution juvenile; and (2) the group (SVP3C1, GC3735, SVP4A) have $\epsilon_{\text{Nd}}(t)$ values that are substantially more negative, between -3.4 and -5.5, indicating increased participation of crustal material.

Figure 5. Images transmitted light microscopy images of zircon and concordia diagram rhyolite sample SVP3C showing a crystallization age of 1882 ± 11 Ma.

Table 3. LA-MC-ICPMS U-Pb data for sample SVP3C

Zircon crystals	Isotope ratios								Age (Ma)				Rho	Th/U	Conc. (%)
	$^{206}\text{Pb}/^{204}\text{Pb}$	$^{207}\text{Pb}/^{206}\text{Pb}$	1 σ %	$^{207}\text{Pb}/^{238}\text{U}$	1 σ %	$^{206}\text{Pb}/^{238}\text{U}$	1 σ %	$^{207}\text{Pb}/^{206}\text{Pb}$	1 σ (abs)	$^{207}\text{Pb}/^{235}\text{U}$	1 σ (abs)	$^{206}\text{Pb}/^{238}\text{U}$	1 σ (abs)		
03Z1	201630	0.11539	0.3	5.9164	1.1	0.37187	1.1	1886.0	5.2	1963.6	9.6	2038.2	18.6	0.96	108.07
05Z3	393153	0.11504	0.4	5.5176	0.8	0.34784	0.7	1880.6	7.2	1903.3	6.6	1924.3	10.9	0.82	102.33
07Z5	59199	0.11433	0.4	5.4968	0.7	0.34870	0.6	1869.3	7.5	1900.1	6.1	1928.4	9.7	0.76	103.16
08Z6	117941	0.11331	0.4	6.0893	3.2	0.38975	3.2	1853.2	6.6	1988.7	27.7	2121.7	57.0	0.99	114.48
09Z7	288412	0.11514	0.5	5.3800	0.7	0.33890	0.5	1882.0	9.6	1881.7	6.4	1881.4	8.4	0.61	99.97
10Z8	265204	0.11331	0.5	5.2940	0.7	0.33886	0.6	1853.1	8.3	1867.9	6.3	1881.2	9.4	0.85	101.51
13Z9	178249	0.11544	0.3	5.4353	0.7	0.34148	0.7	1886.7	4.9	1890.4	6.2	1893.8	10.9	0.91	0.30
14Z10	79442	0.11570	0.3	5.1474	0.6	0.32268	0.6	1890.7	4.9	1844.0	5.5	1802.8	9.1	0.87	0.41
16Z12	108987	0.11354	0.8	5.4625	1.3	0.34894	1.0	1856.8	14.3	1894.7	10.9	1929.5	16.5	0.90	0.30
19Z15	26143	0.11689	0.7	5.3369	1.0	0.33113	0.8	1909.2	11.7	1874.8	8.9	1843.8	13.1	0.76	0.31
20Z16	218499	0.11522	0.7	5.4806	1.1	0.34500	0.8	1883.3	12.6	1897.6	9.2	1910.7	13.3	0.87	0.31
24Z18	31612	0.11441	0.3	5.9788	0.7	0.37899	0.6	1870.7	5.2	1972.8	6.2	2071.6	11.5	0.89	0.37
27Z21	51887	0.11609	0.7	5.3878	1.1	0.33660	0.9	1896.9	12.4	1882.9	9.6	1870.3	14.4	0.77	0.38
28Z22	4679	0.11691	0.3	5.8626	1.0	0.36369	0.9	1909.5	6.2	1955.7	8.5	1999.7	15.8	0.93	0.66
30Z24	5236	0.11676	0.8	6.0914	1.2	0.37836	1.0	1907.3	13.7	1989.0	10.7	2068.6	16.9	0.90	0.27
33Z25	24738	0.11570	1.0	5.6184	1.5	0.35218	1.2	1890.9	17.3	1918.9	13.3	1945.0	20.3	0.77	0.26
34Z26	108631	0.11595	0.5	5.9362	0.9	0.37131	0.8	1894.7	8.9	1966.5	8.2	2023.5	14.1	0.83	0.31
35Z27	946051	0.11368	0.4	5.8771	1.0	0.37497	1.0	1859.0	6.8	1957.9	8.9	2052.7	16.8	0.92	0.23
36Z28	187485	0.11559	0.9	5.8598	1.1	0.36767	0.7	1889.1	16.3	1955.3	10.0	2018.4	12.3	0.75	0.23
37Z29	2306	0.11539	0.4	6.1180	1.5	0.38455	1.4	1885.9	7.6	1992.8	13.0	2097.5	25.6	0.96	0.35
39Z31	135266	0.11452	0.4	5.5367	0.8	0.35064	0.6	1872.4	7.8	1906.3	6.6	1903.7	10.6	0.79	0.43

Table 4. Sm-Nd data for the effusive rocks of the Iricoumé Group

Samples	Lithologies	Sm (ppm)	Nd (ppm)	$^{147}\text{Sm}/^{144}\text{Nd}$	$^{143}\text{Nd}/^{144}\text{Nd}$ ($\pm 2\text{SE}$)	$\epsilon_{\text{Nd}}(0)$	$\epsilon_{\text{Nd}}(t=1.882 \text{ Ma})$	$T_{\text{DM}}(\text{Ga})$	$f_{\text{Sm/Nd}}$
SVP3C1	Rhyodacite	10.29	55.06	0.1129	0.511425 ± 7	-23.66	-3.43	2.46	-0.43
GC3735	Rhyolite	5.93	29.60	0.1212	0.511477 ± 7	-22.64	-5.43	2.59	-0.38
C6604	Rhyolite	8.68	55.04	0.0954	0.511385 ± 11	-24.44	0.03	2.16	-0.51
SVP2E	Andesite	5.33	29.65	0.1088	0.511516 ± 11	-21.89	-0.65	2.25	-0.45
SVP4A	Andesite	5.14	28.35	0.1097	0.511370 ± 8	-24.73	-3.74	2.47	-0.44
SVP3A	Andesite	8.00	39.23	0.1234	0.511771 ± 9	-16.92	0.80	2.19	-0.37
SVP3B	Andesite	8.28	38.61	0.1296	0.511770 ± 12	-16.93	-0.73	2.34	-0.34

Figure 6. $\epsilon_{Nd}(t)$ versus time (Ga) diagram applied to acidic and intermediary effusive rocks from Iricoumé Group. Prepared in accordance to depleted mantle model of De Paolo (1981).

These Sm-Nd isotopic data are interpreted as indicative of reworking, during the Orosian, of a previous Rhyacian or Siderian juvenile continental crust (Transamazonian crust crust-forming event), or with probable contamination by Archean crust (Marques *et al.* 2007).

DISCUSSION

The volcanic rocks of the Iricoumé Group present some important geochemical differences especially on aluminum saturation index (A/NK versus A/CNK), K_2O/Na_2O ratio, alkali contents ($Na_2O + K_2O$), as well as Zr + Nb + Ce + Y and contents. On Harker diagrams, the SiO_2 contents clearly indicate the bimodal nature of the magmatism with a compositional gap between intermediate ($SiO_2 < 57.7$ wt%) and acid rocks ($SiO_2 > 67$ wt%), without delineation of geochemical common trends, suggesting different evolution and magmatic sources (Fig. 7). The progressive increase of SiO_2 is followed by decrease in Al_2O_3 , Fe_2O_{3tot} , MgO, CaO, TiO_2 , P_2O_5 and Sr contents, indicating that plagioclase, hornblende, pyroxene, biotite and apatite have an important role in progressive fractional crystallization, especially in intermediate rocks. On the other hand, Na_2O , K_2O and Rb contents tend to show a positive correlation with increase of SiO_2 , suggesting action of the plagioclase and alkali-feldspar fractionation, mainly in acid rocks. Ta, Nb, Zr, Ce and Y behave as incompatible elements with their contents increase towards to acid rocks, probably

likely to alkali-feldspar fractionation or to accumulation of accessory minerals.

The inverse correlation between the less- and more-differentiated rocks on the Rb versus Sr diagram is also noteworthy (Fig. 8), suggesting that plagioclase and alkali-feldspar were the main minerals fractionated during the magmatic evolution. On Hf + La + Ce versus Zr diagram (Fig. 8), note a positive correlation marked by progressive enrichment of HFS elements towards more differentiated rocks, which is the result of fractionation from the accumulation of incompatible elements likely to accessory minerals structure, such as zircon, allanite, apatite and titanite. These petrogenetic characteristic are consistent with the data available for Iricoumé volcanism in the Pitinga mining district (Ferron *et al.* 2010; Pierosan *et al.* 2011a).

Although typical basaltic composition rocks have not been registered in this region, it is possible that some rocks classified here as andesites are originally basalts modified by later hydrothermal process or amygdaloidal texture presence. This is suggested by the contents of some elements that are considered to be unusual for typical andesite, such as variation in MgO, slightly depleted Na_2O , high K_2O , CaO and LOI on whole-rock geochemistry (Tab. 1). Therefore, it is possible that some basaltic rocks had their original chemical composition modified and moved towards the andesitic composition. For further evaluation of crustal contamination, Zr/Y versus Nb/Y and Zr/Y versus Ce/Y ratios of studied andesite samples along with average crust and primordial mantle are plotted (Fig. 8). The samples plot on a trend very close to average crust, which suggests crustal contamination. This pattern associated with negative $\epsilon_{Nd}(t)$ data indicates that these andesites are probably derived either from a modified mantle source, or the original magmas were contaminated with older crustal rocks during ascent and crystallization.

With respect to the tectonic setting of origin, the multicationic R1 versus R2 values diagram indicate that the acid rocks are roughly distributed between late-orogenic and post-orogenic, whereas on the Rb versus Y + Nb diagram, the samples plot on the post-collisional or within-plate domain (Fig. 9A and 9B). This geochemical characteristic is also consistent with the data available for Iricoumé felsic rocks in the Pitinga mining district (Ferron *et al.* 2010; Pierosan *et al.* 2011a). On the other hand, for intermediate rocks tectonic discriminant diagrams applied to basalts, such as Nb^*2 versus $Zr/4$ versus Y ternary (Meschede 1986) and Zr/Y versus Zr (Rollinson 1993), were used. The samples plot in the within-plate basalt fields (Fig. 9C and 9D).

Several geological mechanisms have been suggested to explain the association between A-type granite/rhyolite with basalt in post-collisional or in within-plate tectonic settings (Whalen *et al.* 1996; Creaser *et al.* 1991; Turner

Figure 7. Harker-type diagrams applied to acidic and intermediary effusive rocks from Iricoumé Group (blank circle = acid rocks and filled circle = intermediate rocks).

et al. 1992; Dall'Agnol *et al.* 1994; Bonin 2007). In the southernmost Guyana shield has been recognized that I and A-type magmatism took place between 1.92 and 1.86 Ga in distinct tectonic settings. During the initial stages, the I-type magmatism is represented by the Água Branca Intrusive Suite emplaced in syn- to post-collisional uplift setting (Valério 2006, 2011; Valério *et al.* 2006, 2009). This was followed by a more-evolved stage, represented by the Iricoumé-Mapuera association emplaced in within-plate or post-collisional tectonic settings as volcano-plutonic ring systems associated to an extensional event (Valério *et al.* 2009; Ferron *et al.* 2010; Pierosan *et al.* 2011a). Other authors (e.g. Almeida 2006; Almeida *et al.* 2007), on the other hand, suggest that the emplacement of the Água Branca and Mapuera suites occurred during the same tectonic stage in response to asthenosphere rise producing mantle-derived magmas which promoted extensive re-melting of the lower crust during underplating.

The $\epsilon_{\text{Nd}}(t)$ values presented in this paper suggested mixing of two-magmas sources (mantle- and crustal-derived

magmas) for the Iricoumé volcanism. Pierosan *et al.* (2009) suggest that the slab-breakoff tectonic model could have been involved to explain the participation of two magmatic sources during a post-collisional setting, promoting asthenosphere uplift and leading to the melting of the metasomatized overriding mantle lithosphere (e.g. Davies & Blanckenburg 1995). Another alternative tectonic model would involve mantle plume and lithospheric delamination (e.g. Saunders *et al.* 1992; Elkins-Tanton 2005; Pirajno 2007; Pirajno & Hoatson 2012), but the lack of additional isotopic (Sm-Nd, Sr-Sr, Lu-Hf) and petrological data, as well as seismic tomography investigations, still inhibit the better understanding of the tectonic evolution for the region.

In the southernmost Guyana shield, the extensive Iricoumé volcanism is associated with caldera system complex, identified mainly in the Pitinga mining district (Ferron *et al.* 2002; Pierosan *et al.* 2011b), which displays several eruptive conduits, as well as collapse structures, favouring the larger production of pyroclastic

Figure 8. Rb versus Sr (A), Zr versus Hf + La + Ce (B), Zr/Y versus Nb/Y (C) and Zr/Y versus Ce/Y (D) ratios plots for acidic and intermediary rocks from Iricoumé Group. Primordial mantle of Wood *et al.* (1979) and average continental crust of Rudnick & Fountain (1995) (blank circle = acid rocks and filled circle = intermediate rocks).

Figure 9. Tectonic discriminant diagrams for the Iricoumé Group rocks (blank circle = acid rocks and filled circle = intermediate rocks). (A) Multicationic R1 versus R2 diagram for acid rocks (Batchelor & Bowden 1985); (B) Rb versus Y + Nb diagram for acid rocks (Pearce 1996); (C) Nb*2 versus Zr/4 versus Y triangular diagram for basaltic rocks (Meschede 1986); (D) Z versus Zr/Y diagram for basaltic rocks (Pearce & Norry 1979).

material (Souza *et al.* 2007). On the basis of the study by Bryan (2007) and Bryan & Ernst (2008) proposals, discussed by Klein *et al.* (2012), the high magma volume produced at a short time registered in this region is compatible with Silicic Large Igneous Province (SLIP).

CONCLUSIONS

The Iricoumé volcanism in the southernmost Guyana shield consists of a bimodal association comprising andesite/basalt to rhyolite components. The acid rocks are subalkaline with high-K,

metaluminous-to-peraluminous compositions, geochemically compatible to A-type magmatism emplaced in post-collisional to within-plate tectonic settings. The intermediate rocks are subalkaline with medium- to high- K contents, metaluminous nature and geochemically compared with within-plate and post-collisional uplift toleitic basalts.

The U-Pb age presented here for the rhyolites (1882 ± 11 Ma) is consistent with the Pb-Pb age already reported for the Iricoumé volcanic rocks in this region. The T_{DM} model ages ($2.59 - 2.16$ Ga) and $\epsilon_{Nd}(t)$ ($-5.43 - 0.03$) indicate that the magmatic evolution was preferentially related to reworking of older Paleoproterozoic

(Rhyacian or Siderian) juvenile continental crust with some contribution from mantle-derived basaltic magmas. It is also possible that there was some contamination by Archean crust.

The data presented here supports the within-plate or post-collisional tectonic setting proposed for the Iricoumé volcanism, most likely to crustal uplift and melting promoted by the emplacement of basalt magma in an extensional regime. This extensive volcanic event was associated with caldera system complex with the eventual collapse and magma resurgence during a short-time interval, comparable to other SLIP in other parts of the world.

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