Three series of alkaline dikes are characterized in a restricted area of the Mantiqueira Range: a weakly silica-undersaturated series ([ne < 7], of potassic character, represented by basanite, tephrites, alkali basalts, basaltic trachyandesites, phonotephrites, tephriphonolites, and trachyandesites, an intermediate series ([7 > normative ne > 13], of potassic character, consisting only of basanites and tephrites, and a strongly silica-undersaturated series ([normative ne > 13], of sodic character, consisting of basanites, tephrites, phonotephrites, and phonolites. Ar-Ar data for biotite from a tephriphonolite dike yielded an average age of 85.4 ± 0.4 Ma. All three series show similar evolution trends controlled mainly by early-fractionation of olivine, spinel and clinopyroxene and, secondly, fractionation of plagioclase and apatite. Different degrees of crustal assimilation are suggested for each series. Assimilation processes are responsible for the wide 87Sr/86Sr (0.70447-0.71172) and 143Nd/144Nd (0.51237-0.51244) ranges, for the increased K2O/Na2O, Ba/Sr and Rb/Sr ratios, and for the reduced Ce/Pb ratios. In general, a significant contribution from partially melted host rocks is associated to the weakly silica-undersaturated potassic series. A phlogopite-bearing carbonate-metasomatized heterogeneous peridotite source accounts for the main chemical variations in the least contaminated samples of each series. Geochemical signatures of the strongly sodic alkaline series indicate a greater contribution from carbonated pyroxene-rich veins in the mantle source.

**KEYWORDS:** alkaline magmatism; potassic and sodic series; crustal assimilation; metasomatized mantle source.

**INTRODUCTION**

Cogenetic association of contrasting strongly and weakly alkaline magmatic series is a common feature of many magmatic provinces (e.g., Wilson et al. 1995, Fitton & Upton 1987, Comin-Chiaramonti et al. 1997). Several factors seem to contribute with their differences in magma composition. Small variations in the degree of partial melting of a common source, distinct degrees of crustal assimilation and/or variable physicochemical conditions of crystallization from similar parent melts, and heterogeneous mantle source metasomatism have all been attributed to classical occurrences such as the Hawaiian alkaline volcanism (Clague 1987), the Cantal volcano, the Massif Central (Wilson et al. 1995), the Teide-Pico Viejo volcanic complex, the Canary Islands (Ablay et al. 1998), and the East African Rift System (Furman & Graham 1999).

The Serra do Mar alkaline province (SMAP; Almeida 1983, Thompson et al. 1998, Brotzu et al. 2005), southeastern Brazil, consists of Upper Cretaceous to Paleogene plutonic complexes and dike swarms formed by magmas of varied degrees of evolution from potassic primitive parental
compositions. Brotzu et al. (2005) distinguished two main potassic suites for the dike swarms of the province: a more widespread, strongly silica-undersaturated suite that ranges from ultrabasic to phonolitic rocks, and a weakly silica-undersaturated suite that ranges from basalts to trachytes. Distinct basanite and foidite parental magmas are assigned to the strongly silica-undersaturated suite. Although the evolution of both suites involved fractional crystallization and crustal assimilation, a heterogeneous metasomatized mantle source is believed to be the main control factor in their generation. Despite their regional character, both potassic suites of alkaline magmas are found locally in SMAP. Bennio et al. (2002) described a strongly silica-undersaturated tephriphonolite to phonolite series and a weakly silica-undersaturated to saturated trachyandesite-trachyte series in the Arraial do Cabo Frio peninsula, a restricted area at the far eastern coast of Rio de Janeiro. Scarce local basalites and trachybasalts are suggested as the main parental melts for both series (Bennio et al. 2002).


The Mantiqueira Range consists of Brasilia belt rocks, mainly of the Socorro-Guaxupé nappe, and rocks of the Embu terrane (Vinagre et al. 2014a, 2014b). In Sapucaí Mirim, Santo Antônio do Pinhal and Campos do Jordão, their main outcrops correspond to metasyenogranites, metamonzogranites and biotite gneisses of the Serra da Águia Limpa barrolith (645–630 Ma, Vinagre et al. 2014a) (Fig. 1B). Metasedimentary rocks of the Embu terrane are only found in a restricted sector (Vinagre et al. 2014a), consisting of quartzites with some intercalations of biotite schists. Serra da Águia Limpa rocks indicate intermediate P (5–6 kbar) and high-T metamorphism from amphibolite to granulite facies (Vinagre et al. 2014b). In the Upper Cretaceous, the batholith experienced exhumation to upper crust levels, as indicated by thermochronological studies based on apatite fission tracks (Tello-Saenz et al. 2003).

The Upper Cretaceous alkaline intrusions that occur in the Mantiqueira Range nearby Sapucaí Mirim, Santo Antônio do Pinhal and Campos do Jordão are exclusively related to the Ponte Nova cumulate mafic-ultramafic alkaline massif and its associated dikes (Fig. 1B). In general, most dikes, identified as alkaline lamprophyres, are thought to represent alkaline basalt, basanite, tephrite and nephelinite magmas. However, a complete alkaline series towards phonolite melts is also recognized (Alves et al. 1992, Thompson et al. 1998, Azzone et al. 2009, 2016).

## MAGMATIC CONTEXT / REGIONAL GEOLOGY

Mesozoic to Cenozoic alkaline and alkaline-carbonatitic occurrences are known in the southern-central portion of the Brazilian Platform, being grouped into several provinces by Ulbrich and Gomes (1981), Almeida (1983) and Riccomini et al. (2005). The Upper Cretaceous to Paleocene alkaline intrusions (less than 100 km³ each) and subordinate dikes (Morbidelli et al. 1995, Gomes & Comín-Chiaramoto, 2005) found on the coasts of São Paulo and Rio de Janeiro, and in the Mantiqueira Range fall within the SMAP (cf. Brotzu et al. 2005 and Thompson et al. 1998; Fig. 1A), intruding Neoproterozoic rocks of the Ribeira and Brasilia belts (Heilbron et al. 2004). In these suites, the abundance of evolved types, mostly nepheline syenites and syenites, is remarkable, as reported by several authors (e.g., Bellieni et al. 1990, Brotzu et al. 1992, 1997, 2007, Ulbrich et al. 2005, Gomes et al. 2017, Enrich et al. 2009, Rosa & Ruberti 2017, Melluso et al. 2017), in the São Sebastião, Búzios and Monte de Trigo Islands and Morro de São João, Itatiaia and Poços de Caldas, for example. On the other hand, the Ponte Nova mafic-ultramafic massif is the only known intrusive suite with prevalence of mafic to ultramafic cumulate rocks (Azzone et al. 2009, 2016). Dike swarms associated with the complex usually include more primitive, basic alkaline rocks (Alves et al. 1992, Garda et al. 1995, Valente 1997, Thompson et al. 1998, Brotzu et al. 2005, 2007, Azzone et al. 2009, Enrich et al. 2009).

Several authors report the presence of dikes of alkaline affinity in Campos do Jordão, a region in the Mantiqueira Range (e.g., Melcher & Melcher 1972, Alves 1978, Cavalcante et al. 1979, Alves et al. 1992, Thompson et al. 1998, Azzone et al. 2009, Menezes et al. 2015). The information presented by them regards mainly geologic and petrographic aspects of these occurrences.

In situ mesocratic to melanocratic alkaline dikes outcrop on many sites along SP-123 road, from Campos do Jordão to Santo Antônio do Pinhal, and even the Ponte Nova Massif neighborhood in the city of Sapucaí Mirim.
The dikes, which represent either single magma pulses or multiple intrusions, appear as steeply dipping, 0.2 to 1.0 m thick dark gray to black intrusions. Zonation is present in most cases, with macrocryst-rich cores and a fine-grained matrix that grades to macrocryst-poor rims in an aphanitic matrix. An NE-trending direction prevails for the dikes intruding the Serra da Água Limpa batholith. A NW-trend is prevalent in the bodies that intrude the Ponte Nova massif neighborhood. Although fresh rock outcrops are scarce, unaltered samples could be collected from blocks found in the highlands drainage system, without zonality control.

Leucocratic to hololeucocratic dikes and sills are also present in the Ponte Nova massif neighborhood and, similarly to the mesocratic to melanocratic bodies, they intrude the alkaline body and the Serra da Água Limpa batholith. Apparently, they are more abundant near the massif. In some cases, tabular bodies evolve to irregular brecciated zones rich in country rock fragments. Phonolites, the most evolved

Figure 1. (A) Geological sketch map of the Serra do Mar alkaline province in southeastern Brazil, modified from Thompson et al. (1998). In the gray rectangle, the studied area. (B) Simplified geological map of part of the Mantiqueira Range. Alkaline dikes are represented by black squares (sampling point of fresh rocks) in the main drainage segments and by the strike direction of in situ bodies. PN, Ponte Nova alkaline massif (Azzone et al. 2009); SALB, Serra da Água Limpa batholith; G-SN, gneisses of the Socorro nappe; QS-ET, quartzites and schists of Embu terrane (Precambrian basement units from Vinagre et al. 2014a). R, main roadways; D, drainages. Reference Cities: SAP, Santo Antônio do Pinhal; CJ, Campos do Jordão; SM, Sapucaí Mirim.
rock type present, commonly intrude Precambrian rocks. Rounded xenoliths of syenogranite are irregularly found in the leucocratic dikes and sills, with diffusion rings in an aphanitic matrix, which, at least in a local scale, indicates assimilation processes.

**PETROGRAPHY**

The mesocratic to melanocratic alkaline dikes in the study area are mainly panidiomorphic porphyritic, with a fine-grained to aphanitic matrix, being identified mostly as alkaline lamprophyres (“camptonites”, “monchiquites” and “sannaites”, cf. Le Maître 2002). The coarse- to medium-grained macrocryst population ranges from > 5 to < 20 vol.%, mainly consisting of clinopyroxene and olivine with subordinate kaersutite. Clinopyroxene is the most important macrocryst in most bodies. Several populations of clinopyroxene macrocrysts are represented by hedenbergite, diopside or Ti-augite cores, all exhibiting corrosion textures and oscillatory and/or step zoning. Titanaugite rims are also present. Glomeroporphyritic textures formed mainly by medium- to coarse-grained clinopyroxenes are recognized in some occurrences. Olivine macrocrysts, when not replaced by low-grade deuteric/hydrothermal-related serpentine, carbonates and muscovite aggregates, present normal zoning, with contrasting thin rims lower in forsterite molecules. In some dikes, olivine macrocrysts show gulf corrosion textures with biotite coronas, whereas in others sites they are completely euhedral. Minor inclusions of Chromite and Cr-spinel are found in some olivine macrocrysts. Kaersutite macrocrysts are less common, displaying normal step zoning, sometimes evolving to pargasite compositions. This lamprophyre assemblage cannot simply be said to form phenocrysts, but it could be interpreted mainly as an assemblage of antecrysts and even xenocrysts. Disequilibrium microtextures are representative of complex plumbing systems, as reported for similar occurrences (e.g., Menezes et al. 2015, Ubide et al. 2012, 2014). Their macrocrysts are basically composed of Ti-augite, plagioclase, kaersutite, Ti-magnetite, alkali feldspar, biotite, apatite, and analcime. Very fine-grained elongated alkali feldspar grains and equivalent amounts of interstitial analcime constitute the felsic mesostasis, possibly related to volatile-rich late magmatic or hydrothermal glass alteration. Ameboid- to globular-shape structures (ocelli) disseminate through the lamprophyres, consisting mainly of carbonates and/or analcime. Rounded centimeter xenoliths of the metasyenogranitic host rock are occasionally found. Some dikes are mineralogically similar to lamprophyres. However, in addition to a coarser matrix, they also contain minor amounts of plagioclase in the macrocryst population. Petrographically, these rock types, which are identified as tephrites, could represent a shortly evolved stage of the lamprophyre family.

Leucocratic to hololeucocratic alkaline dikes and sills are mostly fine-grained to aphanitic in texture, but sometimes porphyritic with medium to coarse-grained macrocrysts, in gray to dark green colors. Phonolites are hololeucocratic and typically exhibit porphyritic texture, with alkali feldspar as the most important macrocryst, and subordinate nepheline. Their matrices are predominantly fine-grained to aphanitic, mainly formed by tabular alkali feldspar and euhedral to interstitial nepheline arranged in a trachytic to fluidal texture. Acicular aegirine is the main mafic phase, with minor amounts of amphibole and apatite. Trachytes, tephritic phonolites and phonolitic tephrites are also found in the study area. These rocks are fine- to medium-grained inequigranular, with greater amounts of tabular plagioclase, pargasite and Ti-biotite. They lack aegirine, and their alkaline feldspar, opaque minerals, apatite and nepheline contents are lower than in the phonolites. In some of these intermediate varieties, clusters of coarse-grained plagioclase are also identified, configuring a glomeroporphyritic texture. In other cases, apatite and biotite are medium-grained macrocrysts set in a felsic fine-grained matrix.

**ANALYTICAL METHODS**

Mineral and whole-rock analyses were carried out at geoanalytical facilities in the Institute of Geosciences, University of São Paulo (GeoAnalítica-USP). A Jeol JXA-8600 electron microprobe was used in the analysis of clinopyroxene, olivine, amphibole and biotite, at an acceleration potential of 15 kV, a probe current intensity of 20 nA, and a beam diameter of approximately 5 μm. Silicate minerals and synthetic oxides were employed as standards, data being corrected online using the PROZA software (Bastin et al. 1984). Whole-rock major element and some trace element analyses were performed on fused glass discs and pressed powder pellets according to analytical methods described by Mori et al. (1999) using a Philips PW 2400 X-ray fluorescence spectrometer. Weight loss on ignition (LOI) was evaluated by standard gravimetric techniques (Mori et al. 1999). Whole-rock trace elements and rare-earth elements were analyzed with a Perkin Elmer ELAN 6100DRC ICP-MS, following procedures described by Navarro et al. (2008).

All isotope analyses were carried out at the geochronological research center at University of São Paulo. Whole-rock Sr and Nd isotope composition analyses
were performed by conventional ion exchange chromatography combined with thermal ionization mass spectrometry (TIMS) following the analytical procedures published in Souza (2009) and Petronilho (2009). For $^{40}\text{Ar}/^{39}\text{Ar}$ dating, samples were irradiated for 24 hours using the IEA-R1 nuclear reactor. Isotope analyses were performed by incremental heating with an ARGUS VI (Thermo Scientific) mass spectrometer. Gas extraction and purification were made with a Nd:YVO4 laser connected to an ultrahigh vacuum cleanup line (SAES-GP50). Data reduction, including correction for nuclear and background interferences, mass discrimination, J-value determination and graphical representation were all processed with the ArArCALC software (Koppers 2002). $^{40}\text{Ar}/^{39}\text{Ar}$ ages were measured in relation to the flux monitor standard Fish Canyon (28.01 ± 0.04 Ma; Phillips & Matchan 2013). Plateau ages consider $\geq 50\%$ of $^{39}\text{Ar}$ released over at least three continuous steps overlapping at a 95% confidence level.

RESULTS

Mineral chemistry

The composition of the main mafic minerals (Suppl. Tab. A1), which confirms the alkaline character of all occurrences, is in accordance with the conclusions of Brotzu et al. (2005, 2007) in that mineral chemical trends are very similar to those of various Upper Cretaceous SMAP occurrences. Olivine macrocrysts are Mg-rich (Fo$_{87-73}$), with very low Ni (up to 0.005 apfu) and CaO (up to 0.01 apfu). Clinopyroxenes constitute the chief mafic phase in all bodies and are associated with early (macrocrysts) and late (matrix) magmatic stages. Although displaying well-defined optical zonation, their compositions are mainly expressed as a variation of a diopside-hedenbergite solid solution series (Wo$_{34-14}$En$_{46-25}$Fs$_{1-13}$), with late stages enriched in the Fs member (Fig. 2A). Early-stage compositions are very similar to those of the Ponte Nova massif (Azzone 2008). The evolution trend is similar to the initial part of Passa Quatro massif and amphibole-free rocks of the Búzios Island (Brotzu et al. 1992, 1997, Gomes et al. 2017) (Fig. 2B). Amphiboles represent a trend of kaersuite compositions, grading toward pargasite and magnesian hastingsite end members (cf. Leake et al. 1997) (Fig. 2C) similar to that reported for the Ponte Nova massif (Azzone 2008). All biotite crystals belong to the phlogopite-annite series (Tischendorf et al. 2007), and the main compositions are concentrated in the Ponte Nova field (Fig. 2D, Azzone 2008).

Geochemistry

Different alkaline dike series are distinguished in the Mantiqueira Range (Fig. 3A). Although the petrographic character of the most primitive types does not allow for the recognition of clear patterns or mineral sequences, whole-rock geochemical data combined with data on intermediate and more evolved types suggest that three magmatic series could be characterized in the region based on normative nepheline content (Suppl. Tabs. A2-A4). A weakly silica-undersaturated series (WAS; normative ne $< 7$) of potassic character is represented by basanite, tephrites, alkaline basalts, basaltic trachyandesites, phonotephrites, tephriphonolites, and trachyandesites (Suppl. Tab. A2). A potassic intermediate series (IS; 7 $>$ normative ne $> 13$) consists only of basanites and tephritics (Suppl. Tab. A3). A sodic, strongly silica-undersaturated series (SAS; normative ne $> 13$) consists of basanites, tephritics, phonotephrites, and phonolites (Suppl. Tabs. A4 and A5). Dikes cutting the Ponte Nova massif are mainly associated with the WAS series, whereas dikes intruding the Precambrian basement near Campos do Jordão belong to the SAS series. Similar compositional trends were found for the aforementioned series. The mineralogical control of the bulk-chemical composition of the three series and their variable SiO$_2$-undersaturation degrees are highlighted in a R$_1$-R$_2$ diagram (Fig. 3C). All series are basically controlled by early fractionation of clinopyroxene and olivine in a similar way as observed near the Ponte Nova massif, with significant amounts of cumulate rocks (Azzone et al. 2016). The series described in the region by Thompson et al. (1998) are not considered representative of the compositional trends discussed in this study. Phonolites are representative of extremely fractionated peralkaline magmas (A.I. $> 1.0$, Suppl. Tab. A5).

An almost analogous pattern of incompatible element enrichment is presented in WAS, IS and SAS (phonolites excluded) diagrams (Figs. 4A to 4D). However, whole-rock compositions for SAS diagrams are characterized by well-defined positive anomalies of Ba, Sr and well-defined negative anomalies of K, Pb, Ti, Zr and Hf in primitive-mantle normalized spidergrams (Fig. 4C). The IS and WAS series, instead, display progressive enrichments in Rb, K, Pb, U and Th (Figs. 4A and 4B). Due to their extremely fractionated compositions, phonolites exhibit positive anomalies of Nb, Zr, Hf, Pb and Rb and negative anomalies of Ba, Sr and Ti (Fig. 4D).

The distribution chondrite-normalized REE values shows a relatively linear behavior for all three series (Fig. 5), featuring a more conspicuous enrichment in LREE (about 100-600 times) than in HREE (about 10-50 times). Also, REE contents are not much different, except for the higher enrichment of the most primitive SAS. Phonolites, however,
show greater LREE enrichment and higher depletion in intermediate REE, in an upward concave-shape pattern with HREE levels more enriched than those of intermediate-REE.

**Sr and Nd Isotopes**

Whole-rock initial isotopic ratios ($^{87}\text{Sr}/^{86}\text{Sr}_i$ and $^{143}\text{Nd}/^{144}\text{Nd}_i$) for each series of the Mantiqueira Range are listed in Supplementary Tables A2-A5. Values for each series fall within the 0.704471–0.707389 interval for $^{87}\text{Sr}/^{86}\text{Sr}_i$. For $^{143}\text{Nd}/^{144}\text{Nd}_i$, they fall within 0.512260–0.512508, excluding the phonolites. The main trend presents a negative correlation in a $^{87}\text{Sr}/^{86}\text{Sr}_i$ vs. $^{143}\text{Nd}/^{144}\text{Nd}_i$ diagram (Fig. 6A).

Phonolites show very high Rb and low Sr contents, with very high $^{87}\text{Sr}/^{86}\text{Sr}_i$ and a wide variation in $^{87}\text{Sr}/^{86}\text{Sr}_i$ ratios (0.705675-0.711715). Two groups can be defined: one, from $^{87}\text{Sr}/^{86}\text{Sr}_i$ data, follows the main series interval (0.705675-0.705983), and the other one shows higher radiogenic values (0.710482-0.711715). In contrast, $^{143}\text{Nd}/^{144}\text{Nd}_i$ values of phonolites (0.512377-0.512443) fall within the same interval for all series.

**Figure 2.** Mineral compositions of weakly to strongly silica-undersaturated alkaline dike series of the Mantiqueira Range. (A) Clinopyroxene compositions in the Wo-En-Fs system (cf. Morimoto *et al.* 1988). (B) Crystallization trends of clinopyroxenes compared with the main evolutionary trends for clinopyroxenes of alkaline complexes from southeastern Brazil. $\text{Fe}^* = (\text{Fe}^{2+} + \text{Mn} + \text{Fe}^{3+} - \text{Na})$. References: 1. São Sebastião Island (Bellieni *et al.* 1990); 2. Vitória Island (Motoki 1986); 3. Monte de Trigo Island (Enrich 2005); 4. Passa Quatro (Brotzu *et al.* 1992) and Itaitiaia (Brotzu *et al.* 1997); 5. Morro Redondo (Brotzu *et al.* 1989); 6. Morro de São João (Brotzu *et al.* 2007); 7(a, b). Búzios Island (Gomes *et al.* 2017). (C) Amphibole compositions in the Ti of the C-site vs. Mg$_{\#_{\text{amp}}}$ (Mg$_{\#}$/(Mg$_{\#}$+Fe$_{\#}$) in apfu). (D) Biotite compositions on the Al$_{\#}$ vs. Fe/(Fe+Mg) (apfu) diagram. Symbols: strongly silica-undersaturated series, black diamonds; intermediate series, gray squares; weakly silica-undersaturated series, white triangles. Gray fields are representative of the respective mineral compositions in the Ponte Nova massif (Azzone 2008).
In general, the most primitive samples of SAS and IS yield the more isotopically primitive values (i.e. lower \(^{87}\text{Sr} / {^{86}}\text{Sr}\) and higher \(^{143}\text{Nd} / {^{144}}\text{Nd}\)) than WAS. However, these intervals are more isotopically evolved than those determined by Thompson et al. (1998) for the least contaminated basic dikes of the area. Exceptions are two samples, one from SAS and one from IS, both lying within a field similar to that of Ponte Nova intrusions, with low degree of contamination (Fig. 6B). Host rocks are also negatively correlated in a \(^{87}\text{Sr} / {^{86}}\text{Sr}\) vs. \(^{143}\text{Nd} / {^{144}}\text{Nd}\) diagram. Their values are more isotopically evolved than those of the alkaline series (Azzone et al. 2016; Fig. 6A).

**Ar/Ar ages**

\(^{40}\text{Ar} / {^{39}}\text{Ar}\) ages were obtained for three ~1.5-mm-wide biotite macrocrystals from the ST-04H tephriphonolite dike. All crystals were analyzed in high-resolution incremental heating experiments, in ten or more steps up to total fusion at 3W or 5W of laser power. Heating spectra are shown in Figure 7. Corresponding data is presented as supplementary material (Suppl. Tab. A6). Biotite crystals provided very consistent plateau ages of 85.7±0.7 Ma, 85.2±0.6 Ma and 84.7±0.6 Ma, all defined by >90% of \(^{39}\text{Ar}\) released.

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**Figure 3.** Major-element discrimination diagrams for the weakly to strongly silica-undersaturated alkaline dike series of the Mantiqueira Range. (A) TAS (SiO\(_2\) vs. Na\(_2\)O+K\(_2\)O, in wt%) diagram (cf. Le Maitre 2002). Dashed line separates alkaline and subalkaline fields (after Irvine & Baragar 1971). (B) Normative mafics vs. normative nepheline diagram. (C) R\(_1\)-R\(_2\) diagram (cf. De La Roche et al. 1980). (D) Highly-potassic, potassic and sodic series diagram (after Middlemost 1975). Symbols: strongly-alkaline dike series (SAS), black diamonds; intermediate dike series (IS), gray squares; weakly-alkaline dike series (WAS), white triangles; phonolites, black circles. Average mineral poles compiled from Azzone (2008) are representative of the main assemblage found in the Ponte Nova massif. Abbreviations: ap, apatite; bt, biotite; cpx, clinopyroxene; fsp, feldspar; ilm, ilmenite; krs, kaersutite; mag, magnetite; nph, nepheline; ol, olivine; pl, plagioclase. The DT field is representative of primitive dikes described by Thompson et al. (1998). The SALB field is representative of heterogeneous host rocks of the Serra da Água Limpa batholith (cf. Vinagre et al. 2014a). PN1 and PN2 fields are representative of mafic cumulates and non-cumulatic intrusions from the nearby Ponte Nova massif, respectively (Azzone et al. 2016).
Intrusion timing of the alkaline dike series

Thompson et al. (1998) determined the range of emplacement/crystallization ages of the SMAP magmatic rocks as between approximately 80 and 50 Ma. However, new high-quality geochronological data extends the upper limit of that age interval. Sato et al. (2008) presented 84-85 Ma U-Pb ages for zircon and baddeleyite concentrates from...
São Sebastião island. Enrich et al. (2009), based on Ar-Ar biotite ages, reported an average value of 86.6 Ma for the Monte de Trigo alkaline suite. Azzone et al. (2009) determined an average K-Ar age of 87.6 ± 1.3 Ma for biotite of the Ponte Nova mafic-ultramafic alkaline massif. For rocks of the Mantiqueira Range, an Ar-Ar result of 80.1 Ma in biotite of a mafic dike from Campos do Jordão was assigned by Regelous (1993), listed in Thompson et al. (1998). New biotite Ar-Ar ages in Supplementary Table 1 and Figure 7, that average to 85.4 Ma, point to the confirmation of the chronogroup > 80 Ma for SMAP inland occurrences. Actually, the alkaline complexes in offshore islands of the State of São Paulo and the Ponte Nova massif were grouped by Riccomini et al. (2005) in a new subprovince

Figure 6. (A) Variation of $^{87}$Sr/$^{86}$Sr vs. $^{143}$Nd/$^{144}$Nd for weakly to strongly silica-undersaturated alkaline dike series of the Mantiqueira Range and nearby host rocks. Fields from Trindade Island and Alto Paranaíba Alkaline Province (APAP) compiled from Marques et al. (1999), Gibson et al. (1995) and Araújo et al. (2001). Field for dikes and massifs of the Serra do Mar alkaline province (SMP) compiled from Brotzu et al. (2007) and references therein. Mixing-model curves for the least radiogenic samples representative of the most primitive basic alkaline magmatism of the area (dike from Thompson et al. 1998, and a plug from Azzone et al. 2016; symbols, vertical crosses) to end-members of gneiss and metasyenogranite of the Serra da Água Limpa batholith (curves 1-3, data from Azzone et al. 2016) based on the Albarède (1995) formulae are plotted to evaluate crustal contribution processes (after Azzone et al. 2016). Mixing-model Curve 4 represent a pole of a sample in the Group 1 phonolites (MT-68c) to metasyenogranite end-member, indicating that phonolites could also be affected by crustal contribution. (B) Zoom of (A) diagram. DT field is representative of basic dikes of the Mantiqueira Range as described by Thompson et al. (1998). PN1 and PN2 fields are representative of Ponte Nova intrusions showing lower and higher degrees of crustal contamination, respectively (Azzone et al. 2016).
that may correspond to an area of Cenozoic uplift adjoining the Santos basin. There, the alkaline magmatism seems to be related to the evolution of the adjoining continental margin, favoring the reactivation of ancient weakness zones, particularly NE-SW-trending parallel Proterozoic shear zones (Ulbrich & Gomes 1981, Almeida 1983, 1986, Riccomini et al. 2005). This chronogroup is also consistent with new U-Pb perovskite ages determined by Guarino et al. (2013), which define 80-91 Ma for kimberlites and phlogopite picrites and 78-81 Ma for kamafugites of the Alto Paranaíba alkaline province (APAP). It is noteworthy that such age intervals go against the age progression hypothesis by Thompson et al. (1998) associated with the ESE track of the Trindade mantle plume.

**Fractional crystallization**

The almost similar evolutionary trends observed in the TAS and R1-R2 diagrams (Figs. 3A and 3C) for the weakly-to-strongly-alkaline dike series in the Mantiqueira Range) seem controlled mainly by fractional crystallization (Figs. 3A and 3C). Previous thermodynamic crystallization models achieved through the MELTS algorithm using primitive regional basanite dikes (Menezes et al. 2015, Azzone et al. 2016) are in accordance with the petrographic descriptions. They indicate that the early-crystallized phases are olivine, spinel, and clinopyroxene, whereas apatite and plagioclase represent late fractioning phases. The evolutionary trends of these melts point to a progressive reduction of MgO, TiO₂, and CaO and, subordinately, an increase in SiO₂, Al₂O₃ and alkalis (Suppl. Tabs. A2-A5). These trends also show that basanite melts could have evolved mainly from fractional crystallization, reaching tephrite-phonolite and even more extreme phonolite compositions, as observed in the Búzios Island rocks, for example (Gomes et al. 2017). In fact, based on geochemical compositions, the parental magma of all three alkaline dike series could potentially evolve to phonolite compositions from fractional crystallization of a gabbroic cumulate assemblage. However, the high degree of SiO₂-undersaturation of these magmas seems to link the phonolites with the SAS series. However, a compositional gap in this series is found between 50 and 60 wt% SiO₂ (Figs. 3A and 3B), an interval also referred in literature as the “Daly gap”. This gap can be interpreted as a consequence of the viscosity increase in middle evolution steps that would prevent migration of intermediate magma to deep crustal levels (e.g., Thompson et al. 2001). However, although fractional crystallization accounts for the continuous part of the series, other scenarios could be envisaged for the origin of the phonolites and of the SAS compositional gap. For example, partial melting of basanite rocks (like those of SAS) or even of alkaline gabbroic cumulates possibly emplaced at deeper crustal levels could also produce phonolite magmas with similar isotopic ratios, as found by Legendre et al. (2005) in the Ua Pou Island, French Polynesia, and by Avanzinelli et al. (2004) in the Pantelleria volcano, Italy.

At the primitive extreme of each series, the petrographic and whole-rock geochemical data indicates that the lamprophyre samples (basanite and tephrite compositions) are representative of the combination of a magmatic liquid with accumulation of crystals, as already shown by Menezes et al. (2015). This is also evidenced by the behavior of trace elements, in which higher Cr, Ni and Co and
lower Sr and Ba concentrations of the primitive magmas of each series (Suppl. Tabs. A2-A5) are positively correlated with a higher crystal cargo of the samples. The porphyritic character of the most primitive dikes of each series (with predominance of olivine and clinopyroxene macrocrysts) indicates a complex plumbing system prior to the final stage of emplacement at shallow crustal levels. Mega- and macrocryst populations with typical disequilibrium features are mainly referred to as antecrysts (Menezes et al. 2015), similarly to those studied by Azzone et al. (2016) in Ponte Nova occurrences. These antecryst populations could be representative of crystallization stages in a chamber environment, at lower levels in the same magmatic system.

At the evolved extreme, the upward concave-shape pattern of the REE of the peralkaline phonolites, with high 

1995). For these models, two possibly pristine parental magma signatures were assumed for the least isotopically evolved samples from the Mantiqueira Range: one representing the mafic dikes (Thompson et al. 1998), the other one representing a small inner plug in the Ponte Nova massif (Azzone et al. 2016). The chosen parental poles are representative of biotite gneiss and metabasaltic rocks of the Serra da Água Limpa batholith (Azzone et al. 2016). Samples of the three series are plotted in the range of the mixing lines (Fig. 6B), suggesting initial mixing between enriched-mantle parental melts (with lower 87Sr/86Sr and higher 143Nd/144Nd ratios) and partially melted hosts (having higher 87Sr/86Sr lower 143Nd/144Nd ratios), analogous to the scenario proposed for the nearby Ponte Nova massif by Azzone et al. (2016). An initial process, with high-T mantle-magma still dominated by the liquid phase mix and different degrees of partial melts of heterogeneous crustal rocks, could account for the isotopic differences between the least isotopically evolved samples of each series. In general terms, SAS seems less affected by crustal contamination than WAS (Fig. 6B). Geochemical signatures point to crustal contamination, and are in agreement with the isotopic data. SAS presents negative anomalies of K and Pb in the primitive-mantle spidergrams (Fig. 4C), high Ce/Pb (Figs. 8A and 8B) ratios, low K2O/Na2O (Fig. 3D) and Rb/Sr (Figs. 8A and 8C), whereas WAS presents a decreasing trend in Ce/Pb ratios (Fig. 8A) and a progressive enrichment in K (Fig. 3A) and Pb levels, Rb/Sr and Ba/Sr ratios (Fig. 8C), and higher K2O/Na2O (Fig. 3D). Ce/Pb ratios also present a positive correlation with 143Nd/144Nd ratios (Fig. 6B).

It should be noticed that, due to lower T-gradients between magma and host rocks, high levels of contamination are expected at deeper crustal levels (Reiners et al. 1995, Tegner et al. 2005). Therefore, alkaline mantle-derived
melts could have also assimilated crustal material within a magmatic plumbing system at deeper levels before its final emplacement at upper crust levels. The macrocryst population and their reaction textures indicate that these magmas underwent some lower-level chamber stages, better conditions being then set for assimilation.

As for phonolites, the group which presents higher $^{87}\text{Sr}/^{86}\text{Sr}$ ratios and similar $^{143}\text{Nd}/^{144}\text{Nd}$ for the main trend can also be explained by crustal assimilation from phonolite initial compositions. Mixing-model curves were also drawn to evaluate this group (Curve 4 in Fig. 6A). In this case, initial compositions were assigned to a phonolite sample (MT-68C) that follows the main isotopic trend, with chosen crustal poles representing Serra da Água Limpa batholith metasyenogranite (Azzone et al. 2016). Differently from the first ones, owing to the high...

Figure 8. Primitive mantle-normalized (McDonough & Sun 1995) trace element ratios for the weakly to strongly silica-undersaturated alkaline dike series of the Mantiqueira Range and nearby host rocks. (A) (Rb/Sr)$_N$ vs. (Ce/Pb)$_N$ diagram. The inset presents the entire range of (Rb/Sr)$_N$ obtained for host rocks. (B) (Ce/Pb)$_N$ vs. $^{143}\text{Nd}/^{144}\text{Nd}$ diagram. The inset presents the entire range of (Ce/Pb)$_N$ obtained for host rocks. Possible trends of crustal contribution processes are indicated by the dashed lines. (C) (Ba/Sr)$_N$ vs. (Rb/Sr)$_N$ diagram. Fields for low-contaminated (PN1) and highly-contaminated intrusions of the Ponte Nova massif (PN2) are shown for comparison. Host rock data and biotite compositional controls from Azzone et al. (2016). High levels of Ce/Pb for low levels of Rb/Sr and Ba/Sr are compatible with alkaline mantle-derived melts (Orejana et al. 2008), whereas high levels of Rb/Sr and Ba/Sr for low levels of Ce/Pb are typical of the contribution from partial melts of crustal basement rocks. (C) Same as (B), but plotting only low-degree contaminated dikes from each alkaline series, compared with fertile source ideal composition (primitive mantle values, PM) and with the field for kamafugite/kimberlite compositions of the Alto Paranaiba province. All values of Rb/Sr and Ba/Sr are higher than PM, suggesting a metasomatized mantle source, and within the field for kamafugite/kimberlite compositions.
Rb/Sr ratios found for the phonolites, the mixing curve presents a convex-like shape. Contaminated phonolites, by these models, present lower levels of crustal contribution at this stage. Geological evidence also supports possible crustal contamination for some phonolites. The rounded xenoliths of syenogranites with diffusion rings in the phonolite aphanitic matrix attest to this process, at least at a local scale. Due to the extremely evolved compositions of these peralkaline phonolites, trace element ratios were not too sensitive to crustal contamination, differently from the more primitive samples of each series as indicated above.

Crustal contribution played an important role in the differentiation trend of SAS and, especially, WAS. However, it seems that the initial assimilation process could not be the sole responsible for the generation of WAS parental magmas from SAS parental ones, once the least radiogenic samples of each series present similar $^{87}\text{Sr}/^{86}\text{Sr}$ and $^{143}\text{Nd}/^{144}\text{Nd}$ ratios with the same trace-element anomalies, but with different normative nepheline values (Fig. 3D). Therefore, other parameters need to be considered to discriminate both series, such as mantle heterogeneities.

**Mantle source heterogeneities and the role of carbonate-rich metasomatism**

Thompson et al. (1998) and Brotzu et al. (2005, 2007), based on isotopic and geochemical signatures of less contaminated samples of the Serra do Mar alkaline province, proposed a heterogeneous mantle source for the genesis of potassic magmas. Thompson et al. (1998) assigned the geochemical and isotopic signatures of these magmas to mantle sources representative of a mixture between an OIB-component (similar to the composition of Trindade volcanic island) and a subcontinental lithospheric mantle component that resembled Alto Paranaíba kamafugitic rocks. In their petrological model, the source of the OIB-component is attributed to a convective mantle which, during its uprise, was mixed in variable degrees with different melts from fusible metasomites within the sublithospheric mantle. Brotzu et al. (2005, 2007), in turn, interpreted the same signatures as a result of low partial melting of a shallow-level continental lithospheric mantle subject to varying degrees of metasomatization, such as a phlogopite-bearing spinel-lherzolite with minor amounts of titanite and apatite. In this scenario, kamafugite and kimberlite melts similar to those of the Alto Paranaíba province are thought to have been metasomatic agents in the lithospheric mantle. Similarly, Guarino et al. (2013) attributed the origin of the potassic, ultrapotassic and carbonatitic rocks of the Alto Paranaíba province to a source assemblage made up of metasomatized mica-carbonate garnet lherzolite.

Neither Thompson et al. (1998) nor Brotzu et al. (2005, 2007) reported the occurrence of strongly silica-undersaturated dike series of sodic nature in the Mantiqueira Range or the entire the Serra do Mar province, which is considered to be of potassic affinity. However, potassic and sodic alkaline series in close temporal and spatial association is a typical feature of Lages and Eastern Paraguay occurrences, which are also related to the meso-cenozoic alkaline magmatism of the South American Platform (Traversa et al. 1996, Comin-Chiaramonti et al. 1997, Gibson et al. 1999, 2006). For such situations, a phlogopite-bearing carbonate-metasomatized peridotite source was proposed by Comin-Chiaramonti et al. (1997) and Gibson et al. (2006).

Considering these hypotheses, a possible scenario for the generation of the SAS, IS and WAS series in the Mantiqueira Range could involve vein-plus-wall-rock melting (Foley 1992). In such cases, variations in the degree of alkalinity and of incompatible element (IE) enrichment would also be associated to the participation of modified signatures of the mantle wall-rock, their modal variation (e.g., cpx/gt ratios in a peridotite source), and the composition and melting degree of metasomatic veins. A potassic component of the mantle source, attributed to veins of ultrapotassic melts similar to those of the APAP (Thompson et al. 1998, Brotzu et al. 2005, 2007), could be inferred by trace element ratios and by isotopic signatures. In a $(\text{Rb}/\text{Sr})_w$ vs. $(\text{Ba}/\text{Sr})_w$ diagram (Fig. 8C), SAS, IS and WAS least-contaminated primitive samples exhibit an intermediate composition between an enriched mantle component (PM, McDonough & Sun 1995) and Brazilian kamafugites, phlogopite picrites and kimberlites (KM, Comin-Chiaramonti & Gomes 2005, Brod et al. 2005, Guarino et al. 2013). In the diagram, SAS has a more restricted composition than IS and WAS, showing lower Rb/Sr and Ba/Sr signatures. In a $^{87}\text{Sr}/^{86}\text{Sr}$ vs. Nd/Nd diagram (Fig. 6), the SMAP primitive magmas (including those of the Mantiqueira Range), present an intermediate isotopic composition in a mixing-line, having one pole with a signature similar to that of Trindade sodic melts and one pole of average ultrapotassic kamafugites and kimberlites of the APAP (Thompson et al. 1998, Brotzu et al. 2007). Also, high levels of $\text{La}_w/\text{K}_w$ ratio (> 1) are owed to residual phlogopite in the source (Gibson et al. 1999). For the $\text{La}_w/\text{K}_w$ ratio, WAS and IS display values between 1.0 and 2.0, most around 1.0, whereas SAS values are higher than 2.3 (Suppl. Tabs. A2-A5). These restricted Rb/Sr and Ba/Sr levels and the high La/K values indicate that phlogopite acted as a residual mineral in the genesis of SAS.

In fact, SAS presents well-defined K-, Zr-, Hf- and Ti-negative anomalies in a primitive mantle-normalized spidergram (Fig. 4C). The same behavior is noticed for carbonatites worldwide (Zeng et al. 2010), which suggests a
greater contribution of carbonate-veins in the mantle to its generation. Azzone et al. (2013), Menezes et al. (2015) and Chmyz et al. (2017) attributed a CO₂-rich metasomatized mantle source to different basanitic compositions in equilibrium with cumulates of the Juquiá and Jacupiranga alkaline-carbonatitic complexes, as well as lamprophyric dikes of the Ponta Grossa arch. Chemically, a CO₂-rich mantle metasomatism can increase CaO/Al₂O₃, La/Zr, Zr/Hf and decrease Ti/Eu ratios in the source and, consequently, in partial melts (Yaxley et al. 1991, 1998, Rudnick et al. 1993, Green et al. 2004, Azzone et al. 2013). Mineralogically, the action of carbonate-rich melts and fluids in a peridotitic upper mantle reduces the amount of orthopyroxene, increases the concentration of clinopyroxene, and accounts for the presence of wehrlite assemblages (Green & Wallace 1988, Yaxley et al. 1991, Rudnick et al. 1993, Moore & Wood 1998, Dalton & Wood 1993). The alkaline series of the Mantiqueira Range show CaO/Al₂O₃ ratios (0.53-1.21), similar to experimental data for a carbonated pyroxene-rich source (0.6-0.7; Hirschmann et al. 2003) and a carbonated peridotite source (1.0-11.0; Dasgupta et al. 2007), and ratios similar to lamprophyre dikes associated with the Juquiá and Jacupiranga complexes (0.74-1.72; Beccaluva et al. 1992, Gibson et al. 1999, Menezes et al. 2015, Chmyz et al. 2017). Also, they show higher La/Zr ratios (3.2-6.8) and Zr/Hf values (1.0-1.6) and lower Ti/Eu ratios than primitive mantle values (Ti/Eu₀. 43-0.85). SAS presents higher La/Zr and Zr/Hf values than IS and WAS.

Non-modal batch melting models for trace elements by Azzone et al. (2013) were adapted to evaluate the contribution of CO₂-rich metasomatic fluids and/or melts (producing a clinopyroxene-veined network structure) in the hypothesis of a vein-plus-wall-rock mechanism for SAS, IS and WAS (Fig. 9). A comparison between the alkaline series indicate that they derive from different degrees of CO₂-rich metasomatism, SAS showing higher contribution of carbonated pyroxene-rich veins than IS, WAS and most alkaline dikes of the Serra do Mar province (Fig. 9). SAS, on the other hand, presents similar ranges of Laₙ/Zrₙ ratios than lamprophyre dikes near the Juquiá and Jacupiranga complexes, for which a carbonated pyroxene-rich source is attributed (Azzone et al. 2013, Chmyz et al. 2017). The composition of SAS is also similar to that of the intra-plate alkaline basalts of Shandong (China), from which Zeng et al. (2010) evidenced the same carbonatic fingerprints as in the mantle source.

The alkaline series of the present study have a similar range of Hf/ₜ/Yₙ ratios plotted within the Serra do Mar province dikes field (Fig. 9). The wide variation of Hf/ₜ/Yₙ ratios suggests either a variable degree of partial melting or a garnet peridotite source with different cpx/gt proportions (Azzone et al. 2013, Chmyz et al. 2017). In the case of dikes, modal variations in the modal source seem more plausible. In fact, according to Stracke (2012), small-scale heterogeneities (i.e. kilometer scale) in melting portions of mantle sources are expected to occur due to crust-mantle recycling.

As emphasized by Ruberti et al. (2012), the origin of mantle metasomatism events in the southeastern South American Platform can be attributed to infiltration of fluids or melts related to subduction-driven processes involved in the agglutination of Gondwana in the Neoproterozoic (Heilbron & Machado 2003), or to small-degree melting of asthenospheric mantle fractions in a distensive environment related to the breakup of that supercontinent in the Mesozoic (Almeida 1983, Comin-Chiaramonti & Gomes 2005). Regardless of the metasomatism events that affected the lithospheric mantle, the hypothetical scenario for the Mantiqueira Range points...
to a phlogopite-bearing carbonated-metasomatized peridotite in the Upper Cretaceous. The sodic nature of the strongly alkaline series appears to represent a greater contribution of carbonated pyroxene-rich veins in a heterogeneous mantle source.

CONCLUSIONS

In a restricted area of the SMAP in the Mantiqueira Range, three series of alkaline dikes were characterized: a strongly silica-undersaturated series ($\text{normative } n_e > 13$), of sodic character, represented by basanites, tephrites, phonotephrites and phonolites; an intermediate series ($7 > \text{normative } n_e > 13$), of potassic character, with only basanites and tephrites; and a weakly alkaline series ($\text{normative } n_e < 7$), of potassic character, consisting mostly of basanite, tephrites, alkali basalts, basaltic trachyandesites, phonotephrites, tephriphonolites and trachyandesites. Similar evolution trends controlled mainly by early fractionation of olivine, spinel and clinopyroxene, followed by plagioclase and apatite fractionation are indicated for these series.

The three series exhibit different degrees of crustal assimilation, which contributed to the wide variation in $^{87}\text{Sr}/^{86}\text{Sr}$ and $^{143}\text{Nd}/^{144}\text{Nd}$ isotopic ratios, the increased K$_2$O/Na$_2$O and Ba/Sr and Rb/Sr ratios and the decreased Ce/Pb signatures. In general, a greater influence of host rock partial melts is related to the weakly silica-undersaturated potassic series.

A phlogopite-bearing carbonate-metasomatized heterogeneous peridotite source responds for the main chemical variation in the least contaminated samples of each series. Metasomatic events could have generated a veined structure and modified the chemical and modal composition of the mantle source. Therefore, the high CaO/Al$_2$O$_3$ and La/Zr and the lower Ti/Eu ratios as compared with those of primitive mantle compositions, together with K-, Zr-, HF- and Ti-negative anomalies in the primitive-mantle normalized spidergrams found for the strongly alkaline series of sodic character are thought as representative of a greater contribution of carbonated pyroxene-rich veins to the heterogeneous mantle source.

New Ar-Ar data for biotite of a tephriphonolite dike yielded an average age of $85.4 \pm 0.4$ Ma. Such a value confirms the chronogroup $> 80$ Ma for the inland SMAP occurrences and, together with other analytical results, go against the age progression hypothesis of an ESE migration of the proposed Trindade mantle plume model.

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SUPPLEMENTARY DATA

Supplementary data associated with this article can be found in the online version: Supplementary Table A1, Supplementary Tables A2-A5 and Supplementary Table A6.
Alkaline dike series of the Mantiqueira Range


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