

PETROGENESIS OF ALKALINE-CARBONATITIC COMPLEXES OF CATALÃO AND IPANEMA, IN SOUTHERN BRAZIL

VINCENZA GUARINO

Dipartimento di Scienze della Terra, Università “Federico II”, Via Mezzocannone 8, 80134 Napoli

INTRODUCTION

Mesozoic to Cenozoic alkaline magmatism is widespread throughout southern Brazil. The main areas of alkaline magmatism are: Alto Paranaíba Igneous Province (or APIP) and Ponta Grossa Arch.

The APIP (Fig. 1) is located between the São Francisco Craton and the NE border of the Paraná Basin, and cover an area of $\sim 3,000 \text{ km}^2$ with an estimated volume of $> 15,000 \text{ km}^3$ of strongly alkaline and carbonatitic rocks (mostly kamafugites, kimberlites, lamprophyres and carbonatites) found as lava flows, pyroclastic successions and hypabyssal intrusions (Gibson *et al.*, 1995; Brod *et al.*, 2000; Araújo *et al.*, 2001; Melluso *et al.*, 2008).

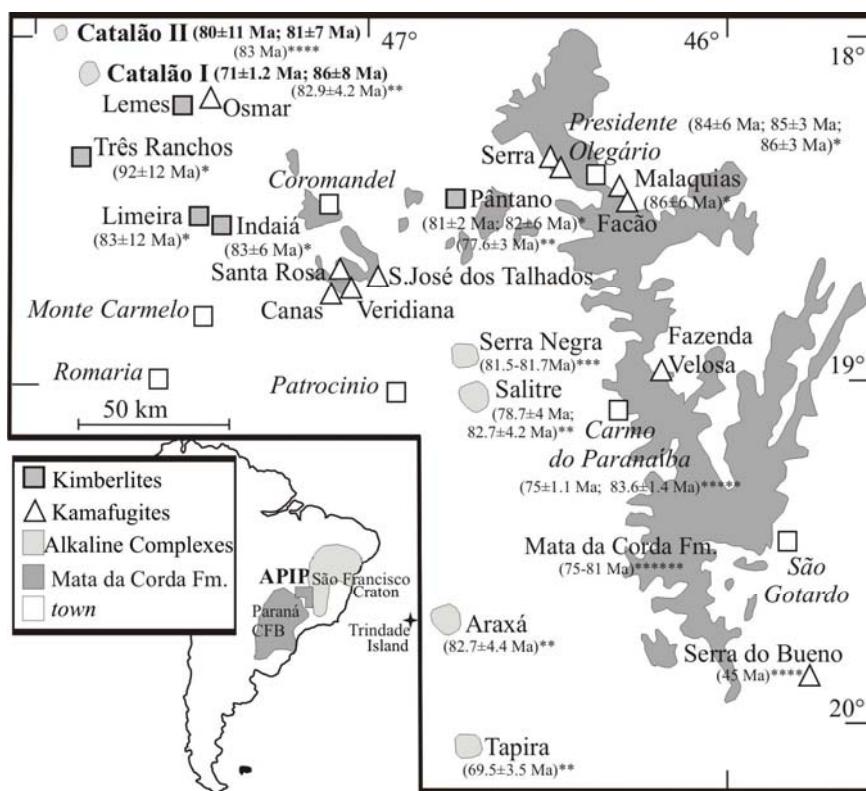


Fig. 1 - Localization of APIP and its main complexes including Catalão I and Catalão II complexes (in bold), southern Brazil (modified after Melluso *et al.*, 2008). Dating are from: (*) Hasui & Cordani (1968); (**) Ulbrich & Gomes (1981); (***) Machado (1991); (****) Gibson *et al.* (1995); (*****) Sgarbi *et al.* (2004).

The Ponta Grossa Arch (Fig. 2) is a large ($\sim 134,000 \text{ km}^2$) tectonic feature on the eastern border of the Palaeozoic-Mesozoic Paraná Basin. This region is characterized by abundant NW-trending tholeiitic dykes. The petrographic associations of alkaline bodies are also quite variable in composition, with the alkaline-carbonatite complexes showing saturated-to-undersaturated syenitic suites related to mafic-ultramafic rocks, nepheline syenites or phonolites.

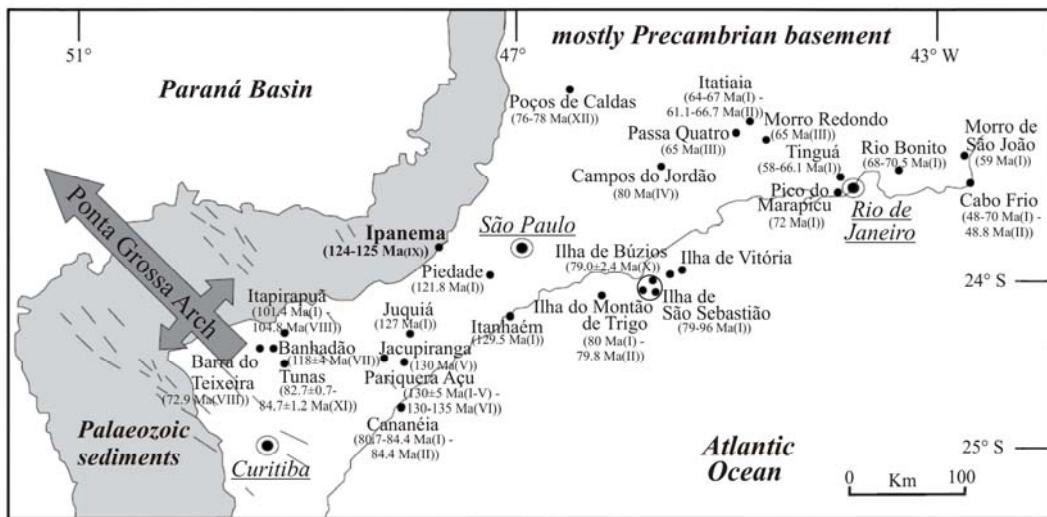


Fig. 2 - Localization of Ponta Grossa Arch and its main complexes including Ipanema complex (in bold). Dating are from: I: Amaral *et al.* (1967); II: Bushee (1974); III: Asmus (1978); IV: Ulbrich & Gomes (1981); V: Amaral (1978); VI: Gomes *et al.* (1990); VII: Hama *et al.* (1977); VIII: Cordani & Hasui (1968); IX: Sonoki & Garda (1988); X: Alves (1997); XI: Siga *et al.* (2007); XII: Shea (1992).

The igneous districts investigated in this paper are: Catalão I and Catalão II (Fig. 1), and Ipanema (Fig. 2), which represent typical alkaline-carbonatite complexes. The aim of this work is to propose a new data set of mineralogical, petrographical and geochemical data. This dataset is used to asses the petrogenetic processes that originated these complexes.

New U-Pb *in-situ* ages are estimated on perovskite, pyrochlore and baddeleyite from Catalão I and Catalão II complexes. The age determinations obtained (from 85 to 71 Ma) are coherent with the previous radiometric investigations (Fig. 1).

Ipanema complex has an age of ~ 125 Ma (Morbidelli *et al.*, 1995).

CLASSIFICATION

Catalão I and Catalão II complexes are composed of ultramafic lamprophyres, carbonatites, phoscorites, fenites, alkaline pyroxenites or clinopyroxenites, apatite-, magnetite- and phlogopite-bearing rocks.

Ipanema complex consists of lamprophyres, carbonatites, shonkinites, feldspar-rich rocks, apatite-, magnetite- and phlogopite-bearing rocks. The carbonatites are classificated as magnesiocarbonatites and calciocarbonatites in Catalão complexes and calciocarbonatites in Ipanema complex, following the chemical classification proposed by Woolley & Kempe (1989). The ultramafic lamprophyres of Catalão

are classified as phlogopite-picrites in according to Gibson *et al.* (1995) and Brod *et al.* (2000), whereas the lamprophyre of Ipanema complex is chemically a tephrite following the chemical classification proposed by Le Bas *et al.* (1986). The rocks called here phoscorites, have been classified on the basis of modal mineral contents, as proposed by Hanekom *et al.* (1965).

The rocks identified as magnetite-, apatite- and phlogopite-bearing rocks are named, respectively, like magnetites, apatites and glimmerites, names used in the literature.

All lithologies observed for these complexes present a holocrystalline granular texture, with cumulitic aspect, except for Catalão phlogopite-picrites with a pseudofluidal porphyric texture and Ipanema lamprophyre with a strongly porphyritic texture.

MINERAL CHEMISTRY

The mineralogical phases, analysed in Catalão I and Catalão II complexes, are:

- carbonate that shows a wide variety in composition, especially in magnesiocarbonatites (dolomite associated with calcite, norsethite, magnesite, stronzianite, alstonite, witherite, burbankite) and in calciocarbonatites (calcite associated with stronzianite, nyerereite, gregoryite, aencylite-Ce, burbankite, siderite, ankerite);

- magnetite is the predominant spinel in Catalão rocks, but it is totally absent in fenites and clinopyroxenites. Ilmenite is present in magnetites, apatites and in calciocarbonatites. Cromites are analysed in phlogopite-picrites, while small crystals of rutile in calciocarbonatites;

- clinopyroxene is present as aegirine-augite in fenites ($\text{Na}_{23-20}\text{Mg}_{62}\text{Fe}^{2+}_{18-15}$), in clinopyroxenites ($\text{Na}_{46-17}\text{Mg}_{59-40}\text{Fe}^{2+}_{24-14}$) and in calciocarbonatites ($\text{Na}_{77-24}\text{Mg}_{48-11}\text{Fe}^{2+}_{28-13}$);

- amphibole, as richterite, is an accessory phase in fenites, clinopyroxenites and rarely in calciocarbonatites;

- olivine is present exclusively in phoscorites ($\text{Fo}_{97}\text{-}\text{Fo}_{98}$) and in phlogopite-picrites ($\text{Fo}_{88}\text{-}\text{Fo}_{89}$);

- alkali-feldspar is present only in fenites as orthoclase ($\text{Or}_{99.97}\text{Ab}_{0.2}\text{An}_{0.1}$);

- garnet is found in phlogopite-picrites (andradite 4-48% and morimotoite 29-68%; Locock, 2008);

- apatite is analysed in any type of rocks, while monazite are present only in magnesiocarbonatites;

- perovskite, pyrochlore and baddeleyite. Perovskite is present only in phlogopite-picrites, shows high CaTiO_3 (88-97 mol.%) contents and variable values in loparite ($\text{Ce}_{0.5}\text{Na}_{0.5}\text{TiO}_3$, 2.2-9.1 mol.%) end-members. Pyrochlore is analysed in phoscorites, magnetites, apatites and rarely in magnesiocarbonatites. Baddeleyite is present in small crystals in apatites and magnetites. Perovskite and pyrochlore are enriched in light lanthanides (La_n/Yb_n , 768.1-872.1 in phlogopite-picrites, and 629.1-842.7 in magnetites and 1411.7 in apatites, respectively), whereas the baddeleyite is relatively depleted in light lanthanides (26.4-33.3 in magnetites and 12.5 in apatites);

- phlogopites belong to phlogopite-tetra-ferriphlogopites series. Chemical and structural analyses display the presence of Fe^{3+} in tetrahedral (${}^{\text{IV}}\text{Fe}^{3+}$) and octahedral (${}^{\text{VI}}\text{Fe}^{3+}$) site. The ${}^{\text{VI}}\text{Fe}^{3+}$ defines the reaction Fe^{3+} -oxy, where excess charge in octahedral site, is compensated in anion site with the addition of OH.

The mineralogical phases, analysed in Ipanema complex, are:

- clinopyroxene is diopside in lamprophyre ($\text{Mg}_{50-76}\text{Na}_{9-20}\text{Fe}^{2+}_{15-30}$) and aegirine-augite in feldspathic rocks ($\text{Mg}_{17-60}\text{Na}_{21-39}\text{Fe}^{2+}_{19-44}$), in shonkinites ($\text{Mg}_{12-44}\text{Na}_{32-64}\text{Fe}^{2+}_{24}$) and in glimmerites ($\text{Mg}_{7-56}\text{Na}_{28-69}\text{Fe}^{2+}_{16-24}$);

- amphibole shows a wide compositional variation in feldspathic rocks (edenite, arfvedsonite and richterite) and in shonkinites (arfvedsonite and richterite), while in glimmerites and in apatites amphiboles are only richterites;
- phlogopite is always present in Ipanema rock. Its compositions indicate that belong to phlogopite-annite series; and in some of these micas take place the Fe^{3+} substitution in place of Al^{3+} ;
- feldspars are principally alkali-feldspar ($\text{An}_{0.2}\text{Ab}_{5.9}\text{Or}_{89.95}$) in shonkinites, and alkali-feldspar ($\text{An}_0\text{Ab}_{9.14}\text{Or}_{86.91}$) and plagioclase ($\text{An}_{0.45}\text{Ab}_{54.94}\text{Or}_{1.6}$) in feldspathic rocks;
- apatites, present in all rocks, show a low TSI contents that indicate an environment of crystallization in equilibrium;
- magnetite is the predominant spinel in these rocks, but it is totally absent in lamprophyre, shonkinites and apatites. Ilmenite is present only in feldspathic rocks;
- carbonate, exclusively as calcite, is present in calciocarbonatites, shonkinites and in some glimmerites.

ISOTOPE GEOCHEMISTRY

Sr and Nd isotopes were estimated on rocks and minerals for Catalão complexes, and only on rocks for Ipanema complex. The Sr and Nd isotopic ratio were age corrected to 82 Ma for Catalão complexes and 125 Ma for Ipanema complex.

In εNd_i vs. εSr_i diagram (Fig. 3), data for the igneous rocks of Catalão fall in the same field of APIP rocks (kimberlites and kamafugites), suggesting comparable magmatic sources. In contrast, Ipanema rocks are markedly different in having lower εNd_i and higher εSr_i . Ipanema rocks show an isotopic composition similar to the Paraguay potassic rocks.

These isotopic differences indicate that the mantle beneath the South American platform is heterogeneous on large scale.

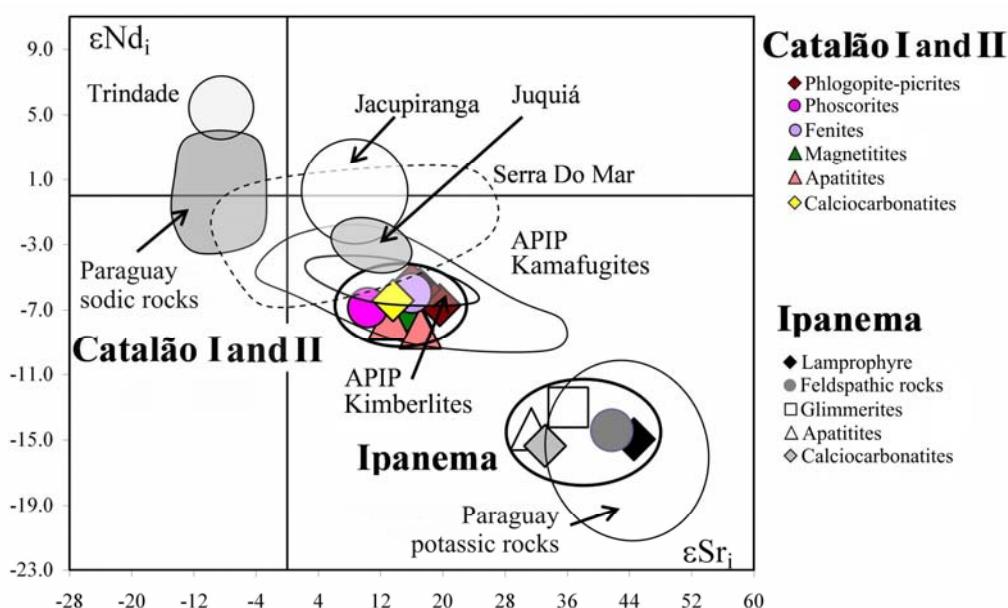


Fig. 3 - εNd_i vs. εSr_i diagram for Catalão and Ipanema rocks.

The O and C isotopic data were obtained on carbonate crystals separated from the Catalão and Ipanema carbonatites. In the $\delta^{13}\text{C}$ (PDB) vs. $\delta^{18}\text{O}$ (SMOW) diagram (Fig. 4), data for the Catalão magnesio- and calciocarbonatites fall within the field of “primary igneous carbonatites”, whereas Ipanema calciocarbonatite fall in “hydrothermal field”.

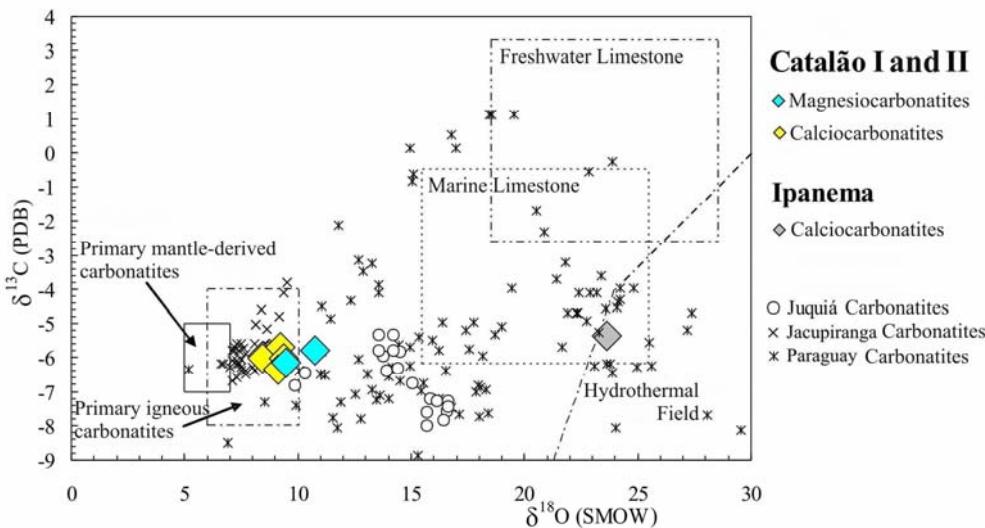


Fig. 4 - $\delta^{13}\text{C}$ (PDB) vs. $\delta^{18}\text{O}$ (SMOW) diagram for Catalão and Ipanema carbonatites.
Data are taken from Comin-Chiaromonti *et al.* (2005) and Castorina *et al.* (1997).

PETROGENESIS OF CATALÃO I AND CATALÃO II COMPLEXES

Source characteristic

Catalão I and Catalão II alkaline-carbonatitic complexes are characterized by phoscorites, magnetitites, apatites, glimmerites, clinopyroxenites, magnesio-, calciocarbonatites and phlogopite-picrites, associated with fenites, which is likely the product of fenitization process following the emplacement of the complexes.

Petrographic, mineralogical and geochemical data show the genetic relationship between the rocks, while the O and C isotopes confirm the magmatic origin of carbonatites.

Phlogopite-picrites could derived by a partial melting of a lherzolitic mantle source enriched in volatiles (H_2O and CO_2) crossed by glimmeritic veins (Foley, 1992a, 1992b).

The SiO_2 undersaturation and the CaO and CO_2 enrichment highlights the lack of orthopyroxene in the source of these magmas, thereby producing a carbonatic metasomatism that converts a lherzolite in a carbonated wehrlite enriched in phlogopite.

Magmatic evolution

Regarding cumulitic rocks are characterised by high K_2O , CaO , Fe_2O_3 , P_2O_5 , H_2O and CO_2 contents and low SiO_2 and Al_2O_3 contents that reflect the peculiar characteristic both of their parental magma and of their different mineralogy.

In the $\text{CaO} - (\text{MgO} + \text{FeO}^*) - (\text{Na}_2\text{O} + \text{K}_2\text{O}) - (\text{SiO}_2 + \text{Al}_2\text{O}_3 + \text{TiO}_2)$ pseudoquaternary system in presence of CO_2 at 1.0 GPa (Lee & Wyllie, 1998; Fig. 5), data for Catalão rocks fall within the

immiscibility fields of carbonatic and silicatic melts, confirming a process of liquid immiscibility for these cumulitic rocks.

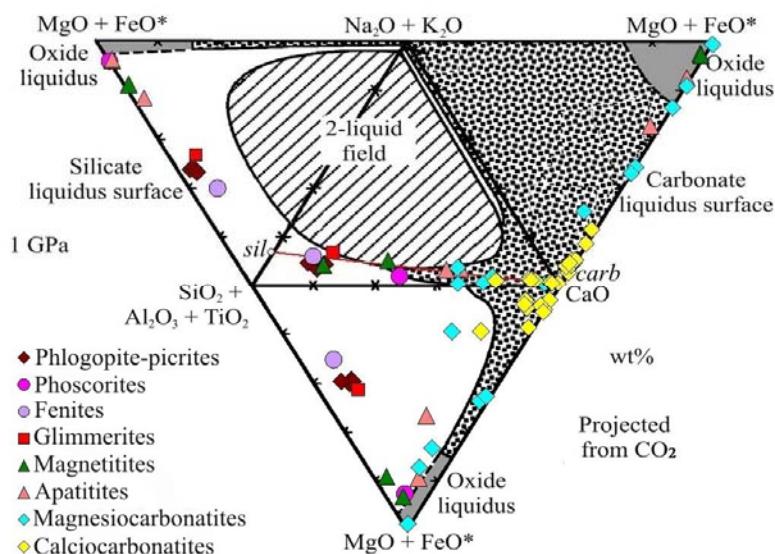


Fig. 5 - Pseudoquaternary system CaO - (MgO + FeO*) - (Na₂O + K₂O) - (SiO₂ + Al₂O₃ + TiO₂) with CO₂, (Lee & Wyllie, 1998) for Catalão rocks.

PETROGENESIS OF IPANEMA COMPLEX

To determine the characteristics source for Ipanema complex is difficult, because lamprophyre, feldspathic rocks and shonkinites are not a parental liquid.

The magmatic evolution of Ipanema rocks can be modelled via fractional crystallization process starting by a parental basanitic magma to produce the cumulitic rocks. The gap in SiO₂ and CaO, around 20-30%, can be a prove of a subsequent immiscibility liquid process that have originated a carbonatitic melts, without significant changes in their geochemical and isotopical features.

In the (SiO₂+Al₂O₃+TiO₂) - (CaO+MgO+FeO*) - (Na₂O+K₂O) system (Lee & Wyllie, 1997; Fig. 6) are reported the AA' and BB' line that represent the comparison of immiscibility fields of carbonate and silicate melts. Ipanema rocks belong to BB' line and albite/calcite (Ab/Cc) cotectic line (Lee & Wyllie, 1998), and both line confirm an evolution trend with initial formation of lamprophyre, feldspathic rocks, shonkinites and glimmerites, and after immiscibility process, are formed apatites, magnetites and calciocarbonatites.

CONCLUSIONS

The main problem of the Catalão I, Catalão II and Ipanema complexes is explain the origin and the close relationship between the alkaline rocks, cumulitic rocks and carbonatites.

Mineralogical, petrographic and geochemical knowledge for Catalão I and Catalão II complexes indicate that the source is a carbonated wehrlite enriched in phlogopite. Carbonatites derived by a liquid immiscibility process. Moreover, the reaction $V^{VI}Fe^{3+}$ -oxy of phlogopites implies a change in the chemical and physical properties within the magmatic chamber.

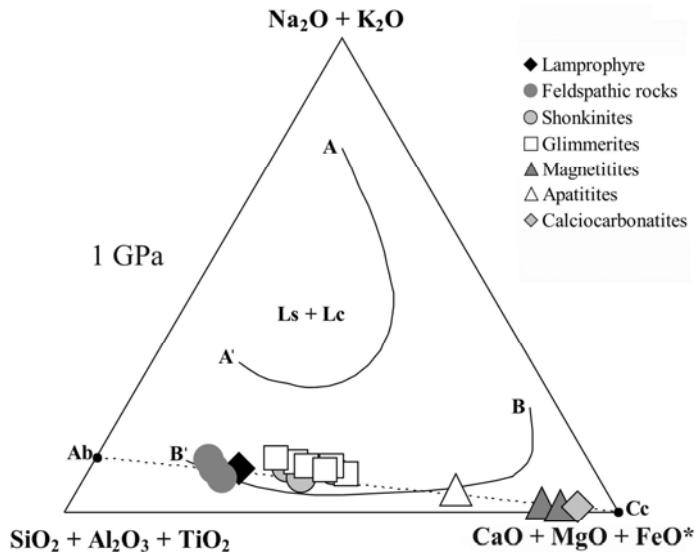


Fig. 6 - $(\text{CaO} + \text{MgO} + \text{FeO}^*) - (\text{Na}_2\text{O} + \text{K}_2\text{O}) - (\text{SiO}_2 + \text{Al}_2\text{O}_3 + \text{TiO}_2)$ with CO_2 system at 1 GPa (Lee & Wyllie, 1997) for Ipanema rocks.

The isotopic evidence shows the close relationship between the Catalão rocks and APIP kimberlites and kamafugites.

The Ipanema rocks derived by fractional crystallization and subsequent liquid immiscibility process, that have originated the calciocarbonatites.

The isotopic compositions show that the Ipanema rocks are much closer to the Paraguay potassic rocks rather than the rocks in the same area, the Ponta Grossa Arch.

REFERENCES

- Alves, F.R. (1997): Contribuição ao conhecimento geológico e petrográfico das rochas alcalinas da Ilha dos Búzios, SP. PhD Thesis, Univ. de São Paulo, 274 p.
- Amaral, G. (1978): Potassium-argon ages studies on the Jacupiranga alkaline district, State of São Paulo, Brasil. Proc. 1st Int. Symp. on Carbonatites, Poços de Caldas, Brasil, 297-302.
- Amaral, G., Bushee, J., Cordani, U.G., Kawashita, K., Reynolds, J.H. (1967): Potassium-argon ages of alkaline rocks from southern Brazil. *Geochim. Cosmochim. Acta*, **31**, 117-142.
- Araújo, A.L.N., Carlson, R.W., Gaspar, J.C., Buzzi, L.A. (2001): Petrology of kamafugites and kimberlites from the Alto Paranaíba Alkaline Province, Minas Gerais, Brazil. *Contrib. Mineral. Petrol.*, **142**, 163-177.
- Asmus, H.E. (1978): Hipóteses sobre a origem dos sistemas de zonas de fratura oceanicas alinhamentos continentais que ocorrem nas regiões sudeste e sul do Brasil. In: "Petrobras. Aspectos estruturais da margem continental leste e sudeste do Brasil", CENPES/DINTEP, Rio de Janeiro (Série Projeto Remac. no. 4), 39-73.
- Brod, J.A., Gibson, S.A., Thompson, R.N., Junqueira-Brod, T.C., Seer, H.J., De Moraes, L.C., Boaventura, G.R. (2000): The kamafugite-carbonatite association in the Alto Paranaíba Igneous Province (APIP) southeastern Brazil. *Rev. Bras. Geocien.*, **30**, 408-412.
- Bushee, J.M. (1974): Potassium-argon ages of some alkaline rocks from southern Brazil. Thesis, Univ. of California, Berkeley, 145 p.
- Castorina, F., Censi, P., Comin-Chiaromonti, P.A., Gomes, C.B., Piccirillo, E.M., Alcover Neto, A., Almeida, R.T., Spezziale, S., Toledo, M. (1997): Geochemistry of carbonatites from Eastern Paraguay and genetic relationships with potassic magmatism: C, O, Sr and Nd isotopes. *Mineral. Petrol.*, **61**, 237-260.

- Comin-Chiaromonti, P., Gomes, C.B., Censi, P., Speziale, S. (2005): Carbonatites from southeastern Brazil: a model for the carbon and oxygen isotope variations. In: "Mesozoic to Cenozoic alkaline magmatism in the Brazilian platform", P. Comin-Chiaromonti & C.B. Gomes, eds. FAPESP, São Paulo, 629-649.
- Cordani, U.G. & Hasui, Y. (1968): Idades K-Ar de rochas alcalinas do primeiro planalto do estado do Paraná. *Anais Congr. Bras. Geol.*, **22**, 149-153.
- Foley, S.F. (1992a): Petrological characterization of the source components of potassic magmas: geochemical and experimental constraints. *Lithos*, **28**, 187-204.
- Foley, S.F. (1992b): Vein-plus-wall-rock melting mechanism in the lithosphere and the origin of potassic alkaline magmas. *Lithos*, **28**, 435-453.
- Gibson, A.S., Thompson, R.N., Leonards, O.H., Dickin, A.P., Mitchell, J.B. (1995): The Late Cretaceous impact of the Trindade mantle plume: evidence from large-volume, mafic, potassic magmatism in SE Brazil. *J. Petrol.*, **36**, 189-229.
- Gomes, C.B., Ruberti, E., Morbidelli, L. (1990): Carbonatite complexes from Brazil: A review. *J. South Am. Earth Sci.*, **3**, 51-63.
- Hama, M., Algarte, J.P., Paiva, I.P., Rodrigues, J.C. (1977): Idades K/Ar do macizo alcalino do Banhado e do complexo Bairro da Cruz. Atas do I Simposio de Geologia Regional, Soc. Brasil. de Geologia, Núcleo de São Paulo, 170-178.
- Hanekom, H.J., van Staden, C.M., Smith, P.J., Pike, D.R. (1965): The geology of the Palabora igneous complex. *South Afr. Geol. Surv. Handbook, Mem.*, 54.
- Hasui, Y. & Cordani, U.G. (1968): Idades potassio-argônio de Rochas Eruptivas Mesozoicas do Oeste Mineiro e Sul de Goiás. *Anais Congr. Bras. Geol.*, **22**, 139-143.
- Le Bas, M.J., Le Maitre, R.W., Streckeisen, A., Zanettin, B. (1986): A chemical classification of volcanic rocks based on the total alkali-silica diagram. *J. Petrol.*, **27**, 745-750.
- Lee, W.J. & Wyllie, P.J. (1997): Liquid immiscibility between nepheline and carbonatite from 2.5 to 1.0 GPa compared with mantle melt compositions. *Contrib. Mineral. Petrol.*, **127**, 1-16.
- Lee, W.J. & Wyllie P.J. (1998): Petrogenesis of carbonatite magmas from mantle to crust, constrained by the system CaO - (MgO + FeO*) - (Na₂O + K₂O) - (SiO₂ + Al₂O₃ + TiO₂) - CO₂. *J. Petrol.*, **39**, 495-517.
- Locock, A.J. (2008): An Excel spreadsheet to recast analyses of garnet into end-member components, and a synopsis of the crystal chemistry of natural silicate garnets. *Comput. Geosci.*, **34**, 1769-1780.
- Machado, D.L. (1991): Geologia e aspectos metalogenéticos do complexo alcalino-carbonatítico de Catalão II (GO). Thèse, Univ. Estadual de Campinas (SP), 102 p.
- Melluso, L., Lustrino, M., Ruberti, E., Brotzu, P., Gomes, C.B., Morbidelli, L., Morra, V., Svisero, D.P., D'Amelio, F. (2008): Major- and trace-element composition of olivine, perovskite, clinopyroxene, Cr-Fe-Ti oxides, phlogopite and host kamacites and kimberlites, Alto Paranaíba, Brazil. *Can. Mineral.*, **46**, 19-40.
- Morbidelli, L., Gomes, C.B., Beccaluva, L., Brotzu, P., Conte, A.M., Ruberti, E., Traversa, G. (1995): Mineralogical, petrological and geochemical aspects of alcaline and alcaline-carbonatite associations from Brazil. *Earth Sci. Rev.*, **39**, 135-168.
- Sgarbi, P.B.A., Heaman, L.M., Gaspar, J.C. (2004): U-Pb perovskite ages for Brazilian kamacitic rocks: further support for a temporal link to a mantle plume hotspot track. *J. South Am. Earth Sci.*, **16**, 715-724.
- Shea, M.E. (1992): Isotopic geochemical characterization of selected nepheline syenites and phonolites from the Poços de Caldas alkaline complex, Minas Gerais, Brazil. *J. Geochem. Expl.*, **45**, 173-214.
- Siga, O., Gomes, C.B., Sato, K., Passarelli, C.R. (2007): O Maciço alcalino de Tunas, PR: novos dados geocronológicos. *Geol. USP Sér. Cient., São Paulo*, **7**, 71-80.
- Sonoki, I.K. & Garda, G.M. (1988): Idades K/Ar de rochas alcalinas do Brasil meridional e Paraguai oriental: compilação e adaptação às novas constantes de decaimento. *Bol. Inst. Geocien., Sér. Cient.*, **19**, 63-85.
- Ulbrich, H.H.G.J. & Gomes, C.B. (1981): Alkaline rocks from continental Brazil. *Earth Sci. Rev.*, **17**, 135-154.
- Woolley, A.R. & Kempe, D.R.C. (1989): Carbonatites: nomenclature, average chemical compositions and element distribution. In: "Carbonatites: genesis and evolution". K. Bell, ed. Unwin Hyman, London, 1-14.