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Late-Hercynian post-collisional dyke magmatism in central Calabria (Serre Massif, southern Italy)

Vanessa Romano^{1,*}, Rosolino Cirrincione¹, Patrizia Fiannacca¹, Michele Lustrino^{2, 3} and Annunziata Tranchina¹

¹ Dipartimento di Scienze Geologiche, Università degli Studi di Catania, C.so Italia, 57, 95129 Catania, Italy ² Dipartimento di Scienze della Terra, Università degli Studi di Roma La Sapienza, P.le A. Moro 5, 00185 Roma, Italy ³ Istituto di Geologia Ambientale e Geoingegneria (CNR - IGAG) Roma, Italy *Corresponding author: vanessa.romano@unict.it

Abstract

Widespread late- to post-collisional magmatism occurred in the Calabria-Peloritani Orogen (southern Italy) during the final stages of the Hercynian Orogeny. In the Serre Massif (central Calabria), medium- to high-K calcalkaline andesitic to dacitic-rhyodacitic dykes show typical geochemical features of subduction-related magmas (LILE and LREE enrichment, HFSE depletion, peaks at Rb, Pb and Th). The origin of late- to post-Hercynian calcalkaline rocks is usually interpreted in an extensional post-collisional framework, involving thinning of the continental lithosphere and progressive passive upwelling of the asthenospheric mantle. In such a context, pure mantle, crustal and hybrid melt production likely can occur. The andesitic dykes were produced by partial melting of an enriched mantle source metasomatized by crustal fluids/melts during former subduction and then suffered minor, if any, assimilation of lower crustal metapelites. Most dacite-rhyodacites were likely derived by hybridization in various proportions of crustal and mantle melts, whereas pure crustal metagedimentary sources, and more or less efficient restite unmixing processes, were involved in the generation of the most silica-rich rhyodacites.

Key words: post-collisional dyke magmatism; andesite; dacite-rhyodacite; Serre Massif; Calabria-Peloritani Orogen.

Introduction

During the late Paleozoic to Early Triassic, a felsic-intermediate and minor mafic magmatism extensively affected large areas of the European Hercynian Belt. Examples are reported from the Pyrenean Chain (e.g., Innocenti et al., 1994; Lago et al., 2004), the French Massif Central (e.g., Ledru et al., 2001; Perini et al., 2004), the Bohemian Massif (e.g., Janousek et al., 2000; Ulrych et al., 2006), the Spanish Central System (e.g., Perini et al., 2004; Orejana et al., 2008), the Sardinian-Corsica Domain (e.g., Traversa et al., 2003; Bonin, 2004) as well as from the Calabria-Peloritani Orogen (Rottura et al., 1990, 1993; Cirrincione et al., 1995; Fiannacca et al., 2008).

Two chronologically, geochemically and mineralogically different magmatic episodes are recognized (Cortesogno et al., 1998, 2004a; Bonin et al., 1998; Bonin, 2004; Rottura et al., 1998; Traversa et al., 2003). The first igneous activity - developed mostly during the Late Carboniferous-Early Permian - produced subalkaline low- to high-K calcalkaline magmas (Cortesogno et al., 1998, 2004a; Cassinis et al., 2008), whereas the latter - mostly Late Permian-Early Triassic - produced transitional to Na-alkaline magmas (Bonin, 1989; Bonin et al., 1998; Cortesogno et al., 1998; 2004b).

Petrogenetic models and geodynamic interpretations proposed for these magmatic events are still matter of debate. In particular, one of the main object of diatribe concerns the geodynamic significance of the magmas belonging to the first cycle, typically showing convergent plate margin geochemical features (Lorenz and Nicholls, 1984; Finger and Steyrer, 1990; Dal Piaz and Martin, 1998). Partially in contrast with this geological reconstructions suggest a post-collisional context and a transtensional regime, developed within an overthickened continental crust suffering a gravity collapse of the main chain, associated with the formation of intermontane troughs (Arthaud and Matte, 1977; Ziegler, 1993; Cortesogno et al., 1998, 2004a; Lustrino, 2000). The occurrence of igneous activity with subduction-related geochemical characteristics in post-collisional settings (e.g., after the end of the oceanic lithosphere subduction) is an anomalous feature, recorded also in other younger orogens, like the Alpine Chain (e.g., Rosenberg, 2004; Lustrino et al., 2011, and references therein).

The Late Permian-Early Triassic phase, characterized by basic-intermediate Na-alkaline magmas up to evolved compositions (e.g., hawaiitic to A-type rhyolitic dykes in Sardinia and Corsica; Atzori and Traversa, 1986; Bonin, 1989; Bonin et al., 1998; Traversa et al., 2003), is instead mostly interpreted as a typical intraplate event preluding to continental break-up and subsequent formation of the Neotethys (e.g., Stampfli et al., 2002).

The present paper focuses the attention on the products of the first phase of activity, cropping out diffusely in the Serre Massif (central Calabria). They consist of calcalkaline intermediate-to-felsic dykes that intrude the metamorphic basement as well as late Hercynian granitoids. The aim of this work is to study petrographic, mineral chemical and geochemical features of these dykes in order to develop a petrogenetic model to explain their origin and to give a contribution on the Late Palaeozoic geodynamic evolution of the southern European Hercynian Belt.

Geological background and field features

The studied dykes crop out in four different areas of the central sector of the Calabria-Peloritani Orogen (hereafter CPO; Figure 1). The CPO is an arcuate orogenic belt bounded by two main tectonic lineaments: the Sangineto line to the North, and the Taormina line, to the South (Tortorici, 1982; Figure 1). The nappe system of the CPO includes several tectonic slices of basement rocks and Mesozoic-Cenozoic sedimentary sequences.

Three are the main hypotheses proposed to explain the origin and geodynamic significance of the CPO:

The CPO would represent a fragment of the Alpine orogenic belt, belonging to the western margin of Adria, emplaced north-westward onto Tethyan ophiolites (Late Cretaceous-Paleogene) (Haccard et al., 1972; Alvarez, 1976; Amodio-Morelli et al., 1976) and later overridden on the Apennine domains (Late Oligocene-Early Miocene);



Figure 1. a) Geological-structural sketch-map of the south-western Mediterranean area (from Ortolano et al., 2005, modified). b) Geological sketch map of the Serre Massif (Southern Calabria-Peloritani Orogen) (Graessner et al., 2000, modified) with location of studied dykes. ST: San Todaro; A: Antonimina; VZ: Villaggio Zomaro; F: Foletti Valley.

The CPO would represent a fragment of the southern European continental paleo-margin, thrust and folded south-eastward onto the Adria margin, starting from middle Miocene, contemporaneously with the opening of the Tyrrhenian Sea (Ogniben, 1969; Bouillin et al., 1986; Knott, 1987; Dewey et al., 1989);

The CPO would represent a micro-continent located between two continental margins, European and African, separated from them by two Tethyan branches and later involved in Europe-Adria collision (Doglioni, 1992; Guerrera et al., 1993; Critelli and Le Pera, 1998; Gueguen et al., 1998; Lustrino et al., 2009; Tortorici et al., 2009; Carminati et al., 2010).

The CPO is classically subdivided into a northern and southern sector, separated along the Catanzaro trough (Tortorici, 1982). The main differences between the two sectors are the lack of ophiolitic units in the southern segment and the presence of an originally Europe-verging chain in the northern-western sector in contrast with exclusively Africa-verging thrusts in the southern one.

The studied dykes crop out in the central part and in the south-western termination of the Serre Massif belonging to the southern sector of the CPO. According to Schenk (1980, 1984, 1990), the Serre Massif represents one of the few places in the world where a nearly complete section of continental crust, with an overall thickness of about 30 km, is exposed. The crustal section is made up of (from the bottom to the top):

a) lower crustal metagabbros, felsic and mafic granulites, and metapelitic migmatites (Schenk, 1980, 1984, 1990; Acquafredda et al., 2006);

b) large bodies of late Hercynian metaluminous to weakly peraluminous granodiorites and tonalites, and minor strongly peraluminous granodiorites and granites, forming the Serre batholith, intruded in between the lower and upper crustal metamorphic rocks (Rottura et al., 1990; Caggianelli et al., 2007 and references therein);

c) greenschist to amphibolite facies metasedimentary and minor metavolcanic sequences, forming the uppermost crustal levels (Colonna et al., 1973; Atzori et al., 1977; Acquafredda et al., 1987; Angì et al., 2010).

Two magmatic suites have been recognized in the Serre batholith: a main metaluminous-toweakly-peraluminous calcalkaline group, representing about the 70% of the exposed granitoids, and a less extensive suite of strongly peraluminous rocks (Rottura et al., 1990, 1993; Caggianelli et al., 2007 and references therein). The first group of rocks show a broad compositional range (~48-70 wt. % SiO₂), with tonalites and granodiorites being the dominant rock types. The strongly peraluminous plutonic rocks lack basic to intermediate lithologies (~67-76 wt. % SiO₂) and contain the typical paragenesis of two micas ± Al-silicates. The metaluminousto-weakly-peraluminous granitoids have been interpreted as I-type rocks resulting from the interaction of mantle-derived magmas with lowercrustal melts (Rottura et al., 1990), whereas the strongly peraluminous granitoids have been interpreted either as typical S-type granites, with sedimentary source rocks (D'Amico et al., 1982; Rottura et al., 1990), or as magmas with mixed mantle-crust origin (Rottura et al., 1991, 1993). According to the previous authors, all the plutonic rocks were emplaced in an extensional regime, during late- to post-collisional phases in the frame of the Hercynian Orogeny. Early geochronological data for the timing of Hercynian magmatism in southern CPO, gave ages spanning the Paleozoic-Mesozoic boundary (from $\sim 298 \pm$ 5 to \sim 270 ± 5 Ma; Rb-Sr whole-rock and mineral ages, zircon U-Pb ages; Borsi and Dubois, 1968; Borsi et al., 1976; Schenk, 1980; Del Moro et al., 1982). More recently, strongly peraluminous granites (Cittanova granite) intruding the Serre batholith in its south-western sector, have been dated by ID-TIMS monazite at 303 ± 0.6 Ma (Graessner et al., 2000) falling in the range of ~304-300 Ma obtained for the strongly peraluminous magmatism in the whole CPO (ID-TIMS monazite and xenotime ages and SHRIMP zircon ages; Graessner et al., 2000; Fiannacca et al., 2008).

Different populations of dykes intrude the plutonic rocks of the Serre batholith as well as the surrounding basement rocks. These dykes have been classified as porphyritic rhyolites, aplites, microgranites and dacites, and include also minor mafic types (Colonna et al., 1973; Borsi et al., 1976; Atzori et al., 1981). Very little is still known about their geochemistry, age, as well as their origin and geodynamic scenario of emplacement. This work aims to investigate four groups of intermediate to felsic dykes. Cropping out in central-southern Serre Massif, in the San Todaro, Foletti Valley, Antonimina and Villaggio Zomaro areas (labelled as ST, F, A and VZ in Figure 1, respectively). The ST dykes are all porphyritic and intrude amphibole-biotite-bearing tonalites and biotite granodiorites. They are dark brown, 1 to 10 m-wide dykes, with tabular or lenticular shape. The grey-green F dykes intrude the phyllite basement with discordant relationships and an average width of 6 m. The light grey A dykes intrude two-mica peraluminous granites and range in width from 0.5 to 3 m. Lastly, the grey-green VZ dykes cut the same peraluminous granites cut by the A dykes with a width ranging from 0.5 to 6 m.

Analytical methods

Whole-rock compositions of 43 samples were determined by X-ray fluorescence (XRF) on powder pellets, corrected for matrix effects (Franzini et al., 1975), at the Dipartimento di Scienze Geologiche, University of Catania. XRF analyses were carried out on a Philips PW 2404 spectrometer, equipped with a Rh anticathode, on pressed powder pellets. Volatile content was measured as loss on ignition (L.O.I.) by standard gravimetric method. Trace element concentrations were obtained by ICP-MS at SGS Mineral Services (Toronto, Canada). 56 elements are detected by sodium peroxide fusion that involves the complete dissolution of the sample in a molten flux. Mineral compositions were obtained using a WDS/EDS-equipped CAMECA SX50 electron microprobe at Istituto di Geologia Ambientale e Geoingegnereia (IGAG)-CNR, Rome, with silicates and oxides as standards, and by SEM-EDS analyses at the Dipartimento di Scienze Geologiche, University of Catania using a Tescan Vega LMU scanning electron microscope equipped with an EDAX Neptune XM4-60 micro-analyzer characterized by an ultra-thin Be window. Analyses were performed at 20 kV accelerating voltage and 0.2 nA beam current. Precision of collected data is on the order of 5%.

Petrography and mineral chemistry

All the samples show a porphyritic texture, with a relatively high Porphyritic Index (P.I.) in F samples. Magmatic assemblages are commonly replaced by secondary mineral associations formed during hydrothermal alteration.

Group A dykes have a P.I. of ~10-15%, with abundant clinopyroxene, and much rarer amphibole and plagioclase phenocrysts. The same phases are present also in the microcrystalline groundmass, together with rare K-feldspar microlites. Colorless clinopyroxene phenocrysts are often fractured and disarticulated, and show corroded cores commonly replaced by chlorite. Brown to pale green amphibole phenocrysts and microcrysts are typically replaced by actinolite. Plagioclase phenocrysts and microcrysts are extensively sericitized. Accessory phases are represented by ilmenite, titanite and apatite. Zircons are very rare and tiny. Among the secondary phases, chlorite, sericite and epidote are the most abundant.

Group F dykes show porphyritic texture (P.I. ~20-25%) with clinopyroxene and amphibole phenocrysts set in a fine-grained groundmass made up of plagioclase, K-feldspar, quartz and amphibole. Clinopyroxene phenocrysts are commonly extensively transformed into chlorite and actinolite aggregates. Hornblende is characterized by high relief, marked cleavage and pleochroism ranging from pale yellow to green. Actinolite replacements are often present at the rim of hornblende phenocrysts, more rarely as total pseudomorphosis. Plagioclase is nearly absent among phenocrysts and is confined to groundmass as extensively sericitized and saussuritized laths. Micrographic intergrowths of quartz and Kfeldspar are also present in the groundmass. Accessory phases consist of ilmenite, apatite, titanite and few and tiny zircon grains. Secondary minerals include chlorite, epidote, sericite and calcite.

Group VZ dykes show porphyritic phaneritic ~10-15%) texture (P.I. with K-feldspar phenocrysts in a medium-grained groundmass made up of subhedral to anhedral and radiating quartz, K-feldspar and white mica. K-feldspar phenocrysts have suffered extensive alteration processes that produced opaque and clay minerals. Quartz rims around K-feldspar phenocrysts suggest reaction processes between phenocrysts and melt. Quartz ocelli, ~1.5-2.0 mm in size, commonly occur as large rounded quartz grains with a K-feldspar and quartz coronitic texture. Local embayment suggest corrosion during melt solidification. Accessory phases are rutile, apatite and zircon. Secondary minerals consist of clay minerals, epidote and white mica.

Group ST dykes can be subdivided in two subgroups in function of textural and mineralogical features: group I, characterized by a porphyritic texture, with a fine-grained matrix and white mica-poor secondary assemblage; group II, characterized by a porphyritic to equigranular texture with a fine to medium-grained matrix, by the presence of abundant interstitial and radial white mica and by micrographic and radiating intergrowths of quartz and K-feldspar. The P.I. is very low (<7%), with extensively altered Kfeldspar and plagioclase phenocrysts. Matrix phases consist of plagioclase, K-feldspar, quartz, chloritized biotite and opaque minerals. Among the phenocrysts, former mafic minerals, possibly amphibole and biotite, have been completely replaced by chlorite. Zircon, rutile and apatite occur as accessory phases.

Pyroxene

In group A samples, clinopyroxene is augite $(Wo_{22-48}-En_{37-64}-Fs_{5-36}; Figure 2; Table 1)$ apart from few crystals that fall in the diopside field (Morimoto, 1988). They all have high MgO

(11.0-19.9 wt.%) and CaO contents (10.4-21.3 wt.%); FeO ranges between 2.9 and 13.6 wt.%, with very low TiO₂ and Na₂O contents (0.23-1.05 wt.%; 0.00–1.66 wt.%, respectively). Mg# value [Mg/(Mg+Fe²⁺)] ranges between 0.69 and 0.88. A slight decrease in Mg#, MgO and CaO and increase in Al₂O₃, FeO and TiO₂ contents, is recorded from core to rim, defining a normal compositional zoning.

Various attempts to determine the composition of the pyroxenes of F dykes did not provide reliable data, probably due to difficulty of obtaining clean analyses in the tiny portions of the crystals apparently spared by chlorite/actinolite replacement.



Figure 2. Clinopyroxene microprobe compositions from A dykes in the Ca-Mg-Fe^{*} (Fe^{*} = Fe²⁺+Fe³⁺+Mn) classification diagram (Morimoto, 1988).

Amphibole

Amphiboles found in Serre dykes are all calcic and are represented by tschermakite and Mghornblende in group A samples (according to Leake et al., 1997; Figure 3 and Table 1); the latter composition also occurs in group F samples. Mg# value of the Mg-hornblende ranges from 0.62 to 0.77, and from 0.73 to 0.98 in A and F dykes, respectively. TiO₂ contents ranges between 0.91-2.33 wt. % and 0.00-2.17

Table 1. Representative major element composition of clinopyroxene and amphibole from Serre	dykes.
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			Cli	korydon	kene					Amph	ibole			Ampł	nibole	
Sample				Ala						A 1	8			H	∞	
Analyses	-	7	б	4	5	9	٢	8	1	7	б	4	1	7	б	4
SiO_2	50.39	52.81	53.43	54.50	53.24	54.82	54.42	53.59	45.18	44.72	43.31	42.99	52.63	54.36	55.40	47.52
TiO ₂	0.00	0.00	0.44	0.00	0.43	0.00	0.00	0.53	2.03	2.42	2.63	2.71	0.50	0.42	0.20	0.77
Al_2O_3	3.38	2.43	2.80	2.92	2.37	3.48	2.64	1.34	9.18	9.90	11.90	11.94	2.93	1.88	0.87	5.80
FeO^{a}	9.47	6.87	5.10	5.74	5.18	6.14	6.37	8.50	15.86	14.90	13.81	12.93	11.04	8.33	12.59	14.45
Cr_2O_3	0.00	0.00	0.32	0.00	0.42	0.00	0.00	0.20	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
MnO	0.00	0.00	0.35	0.00	0.34	0.00	0.00	0.35	0.22	0.20	0.20	0.19	0.34	0.26	0.37	0.37
MgO	13.18	14.23	17.48	16.76	17.97	16.24	16.31	16.56	12.76	13.10	13.25	13.55	17.70	18.88	16.10	15.36
CaO	23.57	23.66	19.52	19.58	19.54	19.33	18.88	18.52	11.12	11.31	10.88	11.02	11.47	12.44	12.36	9.94
Na_2O	0.00	0.00	0.54	0.51	0.51	0.00	0.70	0.42	1.98	2.11	2.19	2.17	0.69	0.51	0.35	1.37
K_2O	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.44	0.44	0.50	0.50	0.14	0.13	0.05	0.22
Total	99.99	100.00	99.98	100.01	100.00	100.01	99.32	100.01	98.77	99.10	98.67	98.00	97.44	97.21	98.29	95.80
Mg#	0.78	0.79	0.86	0.84	0.88	0.83	0.82	0.78	0.72	0.73	0.81	0.82	0.77	0.81	0.70	0.71
Cations are	calculate	d on the	basis of	6 O for	clinopyr	oxene an	d 24 (O	, OH, F)	for ampł	nibole						
Si	1.88	1.96	1.95	1.99	1.94	2.01	2.00	1.98	6.53	6.43	6.20	6.19	7.56	7.72	7.91	7.10
Ti	0.00	0.00	0.01	0.00	0.01	0.15	0.00	0.02	0.22	0.26	0.28	0.29	0.05	0.05	0.02	0.09
Al	0.15	0.10	0.13	0.12	0.10	0.00	0.11	0.05	1.56	1.68	2.01	2.03	0.50	0.31	0.15	1.02
Fe	0.17	0.21	0.16	0.18	0.16	0.19	0.20	0.27	1.92	1.79	1.65	1.56	1.33	0.99	1.50	1.81
Cr	0.00	0.00	0.01	0.00	0.01	0.00	0.00	0.01	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Mn	0.00	0.00	0.01	0.00	0.01	0.89	0.00	0.01	0.03	0.02	0.02	0.02	0.04	0.03	0.05	0.05
Mg	0.73	0.79	0.95	0.91	0.97	0.00	0.90	0.91	2.75	2.81	2.83	2.91	3.79	4.00	3.43	3.42
Ca	0.94	0.94	0.76	0.77	0.76	0.76	0.74	0.73	1.72	1.74	1.67	1.70	1.77	1.89	1.89	1.59
Na	0.00	0.00	0.04	0.04	0.04	0.00	0.05	0.03	0.56	0.59	0.61	0.61	0.19	0.14	0.10	0.40
K	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.08	0.08	0.09	0.09	0.03	0.02	0.01	0.04
Sum	4.00	4.00	4.00	4.00	4.00	4.00	4.00	4.00	15.36	15.41	15.37	15.40	15.25	15.16	15.05	15.52

^a Total Fe as FeO



Figure 3. Calcic amphibole microprobe compositions from Serre dykes in Mg/(Mg+Fe²⁺) vs. Si (apfu) classification diagram (Leake et al., 1997). Symbols as in Figure 2

wt. %, Na₂O between 1.69-2.02 wt. % and 0.00-2.16 wt. %, and K₂O between 0.41-0.90 wt.% and 0.00-0.54 wt. %, respectively. Tschermakite TiO₂ contents show a larger variation range (2.42-6.95 wt. %), as well as Na₂O (0.48-2.19 wt. %); K₂O show similar values (0.00-0.63 wt. %). Secondary amphibole has actinolite-Feactinolite (in A andesites) composition.

Feldspars

In group A dykes plagioclase phenocrysts show secondary albitic composition $(An_{0-6};$ Figure 4a; Table 2). Plagioclase of the groundmass mainly consists of albitic and rare unaltered labradoritic grains (An₀₋₆₄). Orthoclase (Or_{86,2-98,0}) is also present as microlites in the matrix. In group F dykes, plagioclase mainly shows albitic composition in both phenocrysts and groundmass; rare oligoclase occurs as microlites in the matrix (An_{0-25} ; Figure 4b; Table 2). Very thin exsolution of K-feldspar (Or₉₁₋₁₀₀- Ab_{0-9}), optically undetectable, sometimes occur in the albite crystals. Pure orthoclase (Or_{100}) is also present in the groundmass. Plagioclase in ST dykes shows albitic composition with very low anorthite content (An_{0-5} ; Figure 4c; Table 2). Potassium component is generally low $(Or_{0.6-2.9})$; only in few crystals, cores with higher Or component occur (Ab_{84-85} - Or_{14-15} ; Table 2). Orthoclase is characterized by the absence or very low Ab contents (Ab_{0-11} - Or_{100-89}). Several attempts to obtain reliable compositional data on

the feldspars from the VZ dykes resulted unsuccessful because of the pervasive alteration affecting these mineral phases in the studied rocks.

White mica

Textural and chemical criteria, as defined by Miller et al. (1981), allowed us to identify as magmatic only the white mica occurring in the ST (group II) samples. Texturally-defined primary muscovite, characterized by a coarser grain size and well-defined shapes, has higher Na and Ti and lower Si and Mg contents than secondary one. On the whole, primary white mica is here characterized by high Al₂O₃



Figure 4. Feldspars compositions from Serre dykes in An-Ab-Or system. a = A samples; b = F samples; ; c = ST samples.

Sample		A1a		A3c		F8		ST1		ST8	
Location	gm	gm	gm	Ph r	Ph r	Ph c	gm	Ph r	Ph c	Ph c	Ph r
SiO ₂	67.55	67.94	67.60	68.14	65.86	65.69	65.38	67.97	68.04	63.62	67.46
TiO ₂	0.16	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Al_2O_3	20.50	20.80	19.62	19.70	22.16	22.51	18.72	19.69	19.36	22.12	19.51
FeO ^a	0.58	0.32	0.16	0.15	0.00	0.00	0.00	0.06	0.19	0.52	0.21
MgO	0.20	0.13	0.66	0.61	0.00	0.00	0.00	0.70	0.75	0.93	0.72
CaO	1.11	1.13	0.26	0.32	1.30	1.41	5.88	0.03	0.14	0.12	0.22
Na ₂ O	9.88	9.68	11.57	10.99	10.67	10.40	10.02	11.45	11.37	10.16	11.50
K ₂ O	0.00	0.00	0.14	0.08	0.00	0.00	0.00	0.11	0.15	2.53	0.38
Total	99.98	100.00	100.01	99.99	99.99	100.01	100.00	100.01	100.00	100.00	100.00
Atomic p	roporti	ons base	d on 32 ()							
Si	11.80	11.83	11.85	11.90	11.54	11.50	11.63	11.88	11.91	11.33	11.84
Ti	0.02	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Al	4.22	4.27	4.05	4.05	4.57	4.64	3.92	4.05	3.99	4.64	4.03
Fe	0.09	0.05	0.02	0.02	0.00	0.00	0.00	0.01	0.03	0.08	0.03
Mg	0.05	0.03	0.17	0.16	0.00	0.00	0.00	0.18	0.20	0.25	0.19
Ca	0.21	0.21	0.05	0.06	0.24	0.26	1.12	0.01	0.03	0.02	0.04
Na	3.35	3.27	3.93	3.72	3.63	3.53	3.46	3.88	3.86	3.51	3.91
Κ	0.00	0.00	0.03	0.02	0.00	0.00	0.00	0.03	0.03	0.58	0.09
Sum	19.73	19.66	20.10	19.94	19.98	19.94	20.13	20.04	20.04	20.39	20.13

Table 2. Representative major element composition of feldspar from Serre dykes.

^a Total Fe is as FeO

Ph: Phenocryst; r: rim; c: core; gm: groundmass.

(average value \sim 30.0 wt.%) and TiO₂ (0.8-0.9 wt.%) contents and K₂O and MgO values ranging between 9.7-10.4 wt.% and 1.3-1.7 wt.%, respectively (Table 3).

Whole-rock geochemistry

Major and trace element contents of representative samples are reported in Table 4. Most of them were affected by extensive post-magmatic alteration, partially responsible for the relatively high LOI contents (1.4–5.2 wt.%) and low CaO contents. The preferential mobility of alkalies compared to aluminium during hydrothermal stages, is likely responsible for the peraluminous character of many of these dykes [Aluminum Saturation Index (ASI) = molar $Al_2O_3/(CaO + K_2O + Na_2O) = 1.56-4.03$],

particularly evident in some alkali-poor VZ dykes (Table 4). In consideration of the widespread alteration, geochemical classification and modelling of the studied dykes has been essentially based on less mobile or immobile elements.

The SiO₂ content of the studied rocks ranges from ~56 to ~71 wt.%. According to the SiO₂ vs. Nb/Y classification diagram (Winchester and Floyd ,1977; Figure 5) group A and F dykes plot in the andesite field, whereas group ST and VZ dykes mostly plot in the dacite-rhyodacite field. The AFM diagram (Irvine and Baragar, 1971; Figure 6) suggests a calcalkaline affinity for all the investigated dykes. K₂O vs. SiO₂ scheme (Peccerillo and Taylor, 1976) indicates a medium- to high-K calcalkaline composition (Figure 7). Only the SiO₂-poor samples straddle

Sample	819	8111	5112
Location	gm	gm	gm
SiO ₂	48.38	47.64	45.55
TiO ₂	0.94	0.85	0.79
Al_2O_3	30.06	29.63	29.5
FeO ^a	3.04	3.41	5.5
MnO	0.04	0.00	0.02
MgO	1.29	1.31	1.7
CaO	0.02	0.04	0.07
Na ₂ O	0.12	0.14	0.13
K ₂ O	10.43	10.32	9.7
F	0.18	0.04	0.18
Total	94.50	93.38	93.14
Cations ca	lculated on	the basis of 24 (OH, F, Cl)
Si	6.55	6.53	6.34
Ti	0.10	0.09	0.08
Al	4.79	4.78	4.84
Fe	0.34	0.39	0.64
Mn	0.01	0.00	0.00
Mg	0.26	0.27	0.35
Ca	0.00	0.01	0.01
Na	0.03	0.04	0.04
Κ	1.80	1.80	1.72
Sum	13.87	13.91	14.03

Table 3. Representative major element composition of muscovite of Serre dykes.

^a Total Fe as FeO

the line dividing the tholeiitic and the medium-K calcalkaline fields. In these diagrams the composition of late- to post-Hercynian andesitic to rhyolitic calcalkaline dykes from Sardinia-Corsica Domain (SCD) is reported for comparison (Atzori and Traversa, 1986; Atzori et al., 2000; Traversa et al., 2003). A nearly complete overlap between the roughly coeval Serre and SCD dykes clearly emerges.

The group A and F and esites show a relatively high Mg#, ranging from 0.64 to 0.80 and from 0.66 to 0.76, respectively. Ni and Cr contents are high only in F samples (average values for Ni = 128 ppm and Cr = 250 ppm). Major and selected trace element variations are shown in Figures 8-



Figure 5. SiO₂ (wt%) - Nb/Y (ppm) diagram (Winchester & Floyd, 1977). White triangles: A dykes; dark grey triangles: F dykes; white rhombs: ST silicapoor dykes; light grey rhombs: ST silica-rich dykes; dark grey rhombs: VZ dykes.



Figure 6. A $(Na_2O+K_2O wt\%)$ - F (FeO_{tot} wt%) - M (MgO wt%) diagram. Partition line is after Irvine and Baragar (1971). White triangles: A dykes; dark grey triangles: F dykes; white rhombs: ST silica-poor dykes; light grey rhombs: ST silica-rich dykes; dark grey rhombs: VZ dykes. Dashed area includes calcalkaline post-Hercynian dykes of SCD (Atzori and Traversa, 1986; Atzori et al., 2000; Traversa et al., 2003).

Sample	Type	SiO_2	TiO_2	Al ₂ O ₃ I	e2O3	MnO	MgO	CaO	Na_2O	K ₂ 0 1	P_2O_5	LOI	Mg# ⊿	ASI	Cr	Ņ	Rb	Y	Zr Nb	Ba	Sr	
Alb	A	60.26	0.45	12.77	4.8	0.13	9.77	2.69	2.76	1.78	0.17	4.41	80.1	1.8	256 1	02	55	9 15	7 3	311	170	
A3d1	А	59.36	0.56	13.14	5.02	0.18	9.58	1.81	2.94	2.41	0.19	4.82	79.1	1.8	289 1	22	36 1	3 16	3 5	100	173	
A3c	А	61.72	0.62	15.35	3.79	0.09	4.8	1.88	4.61	3.38	0.22	3.53	71.5	1.6	60	25 1	19 1	0 12	6 5	515	189	
Ale	A	60.64	0.6	16.76	3.43	0.09	4.15	2.6	3.01	3.92	0.2	4.59	70.6	1.8	16	10 1	64 1	0 18	1 7	384	146	
A3d2	А	59.81	0.57	14.31	4.6	0.14	8.48	1.6	3.25	2.6	0.19	4.44	78.5	1.9	222	85 8	35 1	3 16	5 6	706	173	
Ala	A	59.69	0.5	13.88	5.02	0.15	9.13	2.23	2.3	2.68	0.18	4.23	78.3	1.9	287 1	03 1	12	0 16	5 5	632	291	
F4	A	59.67	0.52	13.49	6.02	0.12	7.67	2.54	2.96	1.91	0.19	4.91	71.6	1.8	288 1	38	35	0 15	7 5	190	114	
F5	A	60.43	0.52	13.33	5.80	0.12	7.14	2.99	3.27	1.73	0.20	4.47	70.9	1.7	283 1	37 (33	9 14	4	221	184	
F6	A	56.19	0.63	13.63	8.20	0.12	12.89	0.34	2.43	0.61	0.19	4.77	75.7	4.0	299 1	39	00	8 15	8	38	39	
F7	А	57.27	0.61	13.61	7.34	0.12	11.69	0.72	2.77	0.78	0.19	4.92	76.0	3.2	299 1	40	37 1	9 16	1 5	83	75	
F8	A	59.99	0.52	13.12	6.10	0.19	8.25	3.13	3.12	1.38	0.20	4.01	72.9	1.7	290 1	73	12	1 14	.8 33	207	192	
VZ1	D	67.45	0.24	17.38	2.32	0.06	0.76	2.49	0.04	4.61	0.09	4.55	39.5	2.4	7	3 2	48	6 18	8	355	25	
VZ2	D	65.37	0.24	18.24	3.07	0.11	1.48	1.88	0.05	4.60	0.10	4.86	48.8	2.8	б	3 2	42	4 18	7 9	193	16	
VZ4	D	68.33	0.25	20.22	1.82	0.01	0.56	0.26	0.04	5.14	0.11	3.27	37.8	3.7	4	2	60 2	6 19	1 10	175	15	
ST1	D-RD s-p	65.90	0.70	15.20	5.56	0.10	1.31	1.98	4.43	2.97	0.23	1.61	31.9	1.6	lbd	3 1	19 3	6 27	4 11	463	220	
ST3	D-RD s-p	63.43	0.81	15.01	6.92	0.11	3.27	0.79	4.12	2.68	0.23	2.64	48.3	2.0	7	4	11	8 28	9 12	362	147	
ST4	D-RD s-p	67.08	0.63	14.69	5.41	0.09	1.27	1.95	5.47	1.81	0.23	1.38	31.7	1.6	1	4	57	3 26	6 11	377	179	
ST5	D-RD s-p	65.44	0.65	14.81	6.60	0.11	1.77	1.62	4.72	2.33	0.24	1.71	34.7	1.7	б	5	31 4	H 27	1 11	497	169	
ST8	D-RD s-r	69.97	0.20	15.50	3.40	0.02	2.68	0.14	2.89	2.66	0.06	2.47	61.0	2.7	6	4	54 2	9 16	7 9	156	68	
ST9	D-RD s-r	70.97	0.19	14.83	3.24	0.02	2.68	0.11	2.82	2.51	0.05	2.58	62.1	2.7	10	4	46	9 16	0 7	145	63	
ST10	D-RD s-r	70.19	0.22	15.63	3.48	0.03	2.18	0.11	3.20	2.64	0.06	2.27	55.4	2.6	10	3 1	36 2	6 19	5 11	135	67	
ST11	D-RD s-r	67.06	0.38	16.24	4.00	0.04	2.90	0.28	3.46	2.44	0.11	3.09	59.0	2.6	10	5 1	14	5 21	7 7	216	86	
ST12	D-RD s-r	67.02	0.38	16.36	4.42	0.04	2.92	0.25	3.26	2.60	0.10	2.65	56.7	2.7	6	5 1	15 2	14 22	0 7	223	96	
ST13	D-RD s-r	65.73	0.38	16.69	4.69	0.05	3.27	0.29	3.43	2.59	0.11	2.78	58.0	2.6	8	4	16	1 22	L L	235	107	
ST14	D-RD s-r	65.61	0.39	16.46	4.50	0.04	3.67	0.26	3.22	2.69	0.09	3.08	61.8	2.7	11	3 1	14	5 22	6 7	215	85	
ST15	D-RD s-r	66.36	0.39	16.52	4.46	0.04	3.28	0.26	3.19	2.78	0.10	2.63	59.3	2.6	10	5 1	21 2	3 22	L L	212	83	
ST17	D-RD s-r	66.91	0.39	16.53	4.37	0.04	2.86	0.25	3.25	2.62	0.10	2.68	56.5	2.7	10	6 1	16	3 21	6 5	227	79	
ST18	D-RD s-r	68.02	0.37	15.32	4.79	0.06	2.71	0.37	2.88	2.78	0.11	2.60	52.8	2.5	lbdl	4	19 2	14 20	2 6	217	71	
ST25	D-RD s-p	65.71	0.63	14.17	7.00	0.09	3.38	0.51	2.38	2.97	0.18	2.98	48.9	2.4	7	4	11	7 17	0 13	382	73	
ST26	D-RD s-p	65.50	0.66	15.31	6.30	0.08	1.43	1.09	4.00	2.75	0.23	2.67	31.0	2.0	lbdl	2	18 4	1 26	5 11	429	138	
ST27	D-RD s-p	63.85	0.76	15.50	6.49	0.10	2.67	0.69	4.23	2.34	0.25	3.13	45.0	2.1	ŝ	5	4	5 26	6 10	474	122	
ST28	D-RD s-p	62.85	0.81	16.25	6.92	0.12	2.60	0.84	3.53	3.01	0.25	2.83	42.7	2.2	lbdl	5 1	31 4	46 29	0 12	516	203	
ST29	D-RD s-p	64.14	0.73	15.59	5.97	0.11	2.09	1.55	3.61	3.32	0.24	2.67	41.0	1.8	7	5 1	26 3	1 25	7 9	515	196	
ST30	D-RD s-p	64.14	0.72	15.23	6.37	0.11	2.72	1.28	4.34	2.35	0.24	2.51	45.8	1.9	lbd	5	5 20	8 26	0 8	364	147	
A. andes	site. D. Daci	ite. D.RI). darite	-rhvoda	te s-r	v. silica	-2000	r. silica	rich. I	OI = W	eight la	i uo sso	anition									
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Table 4. Major and trace element composition of selected Serre dykes.

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Figure 7. K_2O (wt%) vs SiO₂ (wt%) (Peccerillo and Taylor, 1976). Area as in Figure 6. White triangles: A dykes; dark grey triangles: F dykes; white rhombs: ST silica-poor dykes; light grey rhombs: ST silica-rich dykes; dark grey rhombs: VZ dykes.

9 and reported in Table 4. As a general trend, the dykes show a clear negative correlation with silica only for MgO and Fe₂O₃, whereas the rest of the major oxides show complex or absent correlations. The ST group is characterized by higher Fe₂O₃ compared to the rest of the sample groups for a given SiO₂ content. Another interesting feature is the common distinction of the ST dykes into two main groups, one relatively SiO2-poor (characterized by relatively low MgO and high Fe₂O₃, TiO₂, P₂O₅, CaO and, less clear, Na₂O) and the other relatively SiO₂rich. No substantial differences between the SiO₂-rich and SiO₂-poor ST sub-groups can be observed for K₂O. This geochemical distinction in two sub-groups reflects that already highlighted by the petrographic investigations.

The Serre dykes show complex correlations of trace elements with SiO_2 (Figure 9 and Tables 4, 5). Large Ion Lithophile Elements (LILE) are quite scattered, only Rb showing a slight positive correlation with SiO_2 in A, F and ST samples.

Group F andesitic dykes show positive correlation between LILE and SiO₂, while group A andesites show absent or even negative correlation. The andesites, as a whole group, show lower High Field Strength Element (HFSE; Nb, Hf, Zr) and higher transition element content (Ni, Cr, Sc, V, Co) than dacitic and rhyodacitic dykes. HFSE, Y and Rare Earth Elements (REE) contents in the andesitic dykes are not correlated with SiO₂, differently from what observed for dacites and rhyodacites that show a roughly negative correlation, at least for HFSE. Despite scattering, the observed variation trends are on the whole compatible with the existence of genetic links between samples of the same dyke group, but speak against substantial genetic relationships between the different dyke groups.

Major and trace element compositions of the Serre dykes are compared with those of SCD dykes (Atzori and Traversa, 1986; Atzori et al., 2000; Traversa et al., 2003) and Serre metaluminous to strongly peraluminous granitoids



Figure 8. Major element contents (wt%) vs. SiO₂ (wt%) diagrams. White triangles = A dykes; dark grey triangles = F dykes; white rhombs = ST silica-poor dykes; light grey rhombs = ST silica-rich dykes; dark grey rhombs = VZ dykes. Dashed area: post-Hercynian calcalkaline dykes of the SCD (Atzori and Traversa, 1986; Atzori et al., 2000; Traversa et al., 2003); grey and light grey areas: calcalkaline metaluminous and strongly peraluminous granitoids of Serre Massif, respectively (Rottura et al., 1990).



Figure 9. Selected trace element contents (ppm) vs. SiO_2 (wt%). White triangles = A dykes; dark grey triangles = F dykes; white rhombs = ST silica-poor dykes; light grey rhombs = ST silica-rich dykes; dark grey rhombs = VZ dykes. Dashed area: post-Hercynian calcalkaline dykes of the SCD (Atzori and Traversa, 1986; Atzori et al., 2000; Traversa et al., 2003); grey and light grey areas: calcalkaline metaluminous and strongly peraluminous granitoids of Serre Massif, respectively (Rottura et al., 1990).

Sample	La	Ce	Nd	Sm	Eu	Tb	Yb	Lu	Th	U	Та	Hf
A1b	18.0	40.0	20.4	3.9	1.0	0.5	1.3	0.2	7.1	1.9	0.5	4.0
A3c	18.0	57.0	22.6	4.2	1.2	0.5	1.3	0.2	7.6	2.2	0.5	4.0
Ale	22.0	53.0	21.9	4.2	1.2	0.5	1.3	0.2	7.9	2.1	0.5	4.0
A3d2	24.0	34.0	24.6	4.6	1.2	0.6	1.3	0.2	7.4	2.4	0.5	4.0
Ala	18.0	29.0	20.6	3.9	1.0	0.5	1.3	0.2	7.2	2.0	0.5	4.0
F4	24.6	54.4	21.3	4.2	1.5	0.5	1.3	0.3	7.1	2.0	0.5	4.0
F5	23.1	41.0	20.8	4.2	1.1	0.6	1.4	0.2	7.0	2.1	0.5	3.0
F8	23.2	44.2	23.6	4.9	1.3	0.7	1.8	0.3	7.3	2.3	0.5	3.0
VZ1	52.9	59.8	34.0	6.3	1.0	0.8	2.1	0.3	16.9	4.1	0.7	5.0
VZ2	32.5	59.5	35.0	6.7	1.0	0.9	2.1	0.3	16.8	4.4	0.8	5.0
VZ4	39.4	67.5	38.2	7.0	1.0	0.8	2.2	0.3	18.2	5.0	0.8	6.0
ST4	38.3	81.2	36.9	7.6	1.6	1.0	2.8	0.4	10.8	2.4	0.5	6.0
ST5	39.1	81.1	37.5	7.6	1.7	1.1	3.0	0.5	10.3	2.6	0.5	6.0
ST9	38.3	87.7	35.4	6.7	1.1	1.0	2.6	0.4	14.2	3.1	0.6	5.0
ST10	21.9	50.5	21.7	4.6	0.8	0.8	2.7	0.4	14.4	3.4	0.6	5.0
ST11	18.6	26.1	13.7	3.1	0.7	0.6	2.0	0.4	8.2	2.8	0.5	5.0
ST12	15.9	20.4	10.5	2.5	0.6	0.5	2.0	0.3	8.2	2.9	0.5	5.0
ST13	15.1	16.8	9.7	2.5	0.6	0.5	2.2	0.4	8.6	3.0	0.5	5.0
ST17	12.0	30.4	14.9	3.0	0.6	0.5	1.9	0.3	8.2	2.8	0.5	5.0
ST26	57.6	91.8	46.5	8.6	1.8	1.2	3.1	0.5	11.5	2.9	0.5	6.0
ST27	18.1	35.4	18.9	4.2	0.8	0.7	3.4	0.6	10.7	3.0	0.5	6.0
ST28	28.2	49.9	25.9	5.8	1.1	1.0	3.6	0.6	11.4	3.4	0.6	6.0
ST30	76.8	187.6	67.3	11.3	2.3	1.4	3.2	0.6	11.0	2.8	0.5	6.0

Table 5. Additional trace element analyses by ICPMS of selected Serre dykes.

(Rottura et al., 1990). The greatest differences are seen in the anomalously high MgO and low Al_2O_3 content of the Serre dykes, which are also characterized by low to very low Ca content at any SiO₂ value (Figure 8), all largely reflecting the effects of post-magmatic alteration. Group VZ dykes and SiO₂-rich group ST dykes are also characterized by anomalously low P₂O₅ content. The ST dacitic-rhyodacitic dykes, intruding the metaluminous to weakly peraluminous Serre granitoids, show a rough similarity with these rocks.

Chondrite-normalized REE patterns (Figure 10) are weakly to strongly fractionated, with La_N/Yb_N ratios ranging from 9.9 to 14.0 and from 8.5 to 11.8 for group A and F andesites, respectively. Europium anomaly ranges from slightly negative to slightly positive values (Eu/Eu* = 0.82-1.21). The VZ dacitic samples show REE fractionation similar to that of the andesites ($La_N/Yb_N = 12.9$ -

13.8), but with a more prominent Eu negative anomaly (Eu/Eu* ~0.53). On the other hand, the ST dacitic and rhyodacitic samples show a much more variable REE fractionation ($La_N/Yb_N = 2.6$ -16.3), coupled with clear Eu negative anomaly (Eu/Eu* = 0.53-0.83). The VZ dykes show nearly flat HREE patterns, with Ho_N/Lu_N ratios ranging from 0.95 to 1.71. HREE are nearly constant also in the other dyke groups, clustering around 4-9 times CI chondrite for the andesites, around 10 times for the VZ dacites and around 10-20 times for the ST dacites and rhyodacites.

On the whole, REE element patterns of the Serre dykes share many similarities with the lateto post-tectonic granitoids of the Serre batholith and with the post-Hercynian intermediate to felsic dykes from the SCD (Figure 10).

Primitive mantle-normalized incompatible element patterns (Figure 11) show some common features, i.e., LILE enrichment relative to HFSE,



Figure 10. Chondrite CI-normalized REE patterns for the Serre dykes: a) A samples; b) F samples; c) VZ samples; d) ST samples. Normalization after Nakamura 1974. White area: SCD calcalkaline dykes (Atzori and Traversa, 1986; Atzori et al., 2000; Traversa et al., 2003); grey area: Serre calcalkaline granitoid rocks (Rottura et al., 1990).

peaks at Rb and Pb, negative anomalies at Ba, Nb-Ta, Ti and Sr. Compared with the andesitic rocks, dacitic and rhyodacitic samples show stronger Ba, Sr and Ti negative anomalies and a less pronounced peak at Pb, but overall similar patterns. On the whole, these patterns show typical features of subduction-related magmas (Figure 11; Shimizu and Arculus, 1975; Pearce, 1983; Lustrino et al., 2011).

Discussion

Minero-petrographic and geochemical features suggest, on the whole, an origin from mantle sources modified by fluids produced by metamorphic dehydration reactions during subduction of oceanic slabs for the Serre dykes. The absence of iron enrichment in the first evolutionary stages is probably the strongest evidence for their calcalkaline character (e.g., Arculus, 2003). This, coupled with low TiO₂ and relatively low alkali contents resemble closely the chemical composition of magmas emplaced along active or fossil subduction zones (e.g., Lustrino et al., 2011, and references therein). Unlike typical arc-type rocks, the Serre dykes display a distinct Ba negative anomaly with respect to the adjacent Rb and Th. This trough can be considered not related to the alteration processes but a consequence of crystal fractionation or distinctive source characteristics. This feature also occurs in Plio-Quaternary potassic to ultrapotassic subduction-related rocks from peninsular Italy (Peccerillo and Martinotti, 2006; Avanzinelli et al., 2009; Conticelli et al., 2010, and references therein). Among the incompatible trace elements, the HFSE troughs, the positive Pb anomaly and the high



Figure 11. Primitive mantle normalized trace elements patterns for Serre calcalkaline dykes: a) A and F samples; b) ST and VZ samples. Normalization after McDonough and Sun, 1995. Data field for subductionrelated volcanic is shown for comparison (Gerlach et al., 1988)

LILE/HFSE ratios are classically considered as typical features of subduction-related igneous rocks (Gill, 1981; Grove and Kinzler, 1986; Tatsumi, 1989; Davies and Stevenson, 1992; Hawkesworth et al., 1993; Arculus, 1994; Pearce and Peate, 1995) These characteristics have been commonly explained by the addition of hydrous fluids released from subducting oceanic lithosphere, selectively enriched in LILE and in other fluid-mobile elements (e.g., Pb, U, with high to very high Pb/U ratios), to the mantle wedge, lowering the mantle solidus and leading to magma generation.

The occurrence of such subduction-related geochemical characteristics is at odd with the envisaged tectonic setting of the emplacement of the dykes, being these emplaced after the end of the Gondwana-Laurasia collision (e.g., Stampfli and Borel, 2002). A general agreement exists that late-Hercvnian Calabrian magmas were emplaced in an extensional tectonic setting associated with the post-collisional attenuation of the previously thickened lithosphere in consequence to possible slab detachment (Caggianelli et al., 2007, and references therein). Calcalkaline magma with geochemical features similar to those observed in the intermediate Serre dykes can be produced by decompression melting of an asthenospheric mantle source previously metasomatized by subduction-related fluids/melts (Johnson et al., 1978; Cameron et al., 2003) or of a continental lithospheric mantle previously modified by subduction (Hawkesworth et al., 1995; Wilson et al., 1997). Besides, calcalkaline compositions have been interpreted in literature also as the effect of the interaction between mantle-derived melts with local crust, through wall-rock assimilation and fractional crystallization (Stille and Buletti, 1987; Innocenti et al., 1994; Rottura et al., 1998; Cannic et al., 2002). In all these cases, the presence of calcalkaline magmatism does not automatically imply the presence of coeval subduction. In the next paragraph, we therefore investigate the possible petrogenetic models that may better explain the intermediate-felsic calcalkaline dyke magmatism in the Serre Massif.

Petrogenetic model

In the Th/Yb vs. Ta/Yb discrimination diagram for intermediate-felsic rocks (Pearce, 1983; Gorton and Schandl, 2000; Figure 12a), the Serre andesitic dykes plot in the active continental margin field together with most of the ST silicarich dacite-rhyodacites. The other dacite-rhyodacitic samples straddle the boundary with the volcanic arc field (Figure 12a). Similar considerations can be made using the Rb vs. discrimination diagram for Y+Nb felsic compositions (Pearce et al., 1984; Figure 12b), where the Serre dacites and rhyodacites mainly plot in the Volcanic Arc Granites field (VAG) field, which also includes Active Continental Margin Granites, and near the triple point, as commonly observed for post-collisional granitic rocks (Pearce, 1996).

Dykes with andesitic composition crop out in two localities: Foletti Valley (F group) and Antonimina (A group). The dykes of these two



Figure 12. a) Compositions of Serre dykes in the Th/Yb vs. Ta/Yb plot (Pearce, 1983) modified by Gorton and Schandl (2000) for intermediate-felsic rock compositions. Oceanic Arcs; ACM: Active Continental Margins; WPVZ: Within Plate Volcanic Zones; WPB: Within Plate Basalts. b) Rb (ppm) vs. Y+Nb (ppm) diagram (Pearce et al., 1984) for Serre dacite-rhyodacitic dykes. SYN-COLG: syn-collisional granites; WPG: within-plate granites; VAG: volcanic arc granites; ORG: ocean-ridge granites

areas share many petrographic similarities but can be easily distinguished geochemically, being the group F andesites characterized by much higher Fe₂O_{3tot} (5.40-8.20 wt.%) and lower K₂O (0.61-2.09 wt.%) than group A andesites (3.22-5.02 wt.% and 1.78-4.50 wt.% for Fe₂O_{3tot} and K₂O, respectively; Figure 8). With the exception of the andesites with the lowest SiO₂ content, the group F andesites have also typically higher CaO than group A dykes (Figure 8). On the other hand, the two types of andesites cannot be distinguished on the basis of the compatible and incompatible trace element content (Figures 9, 11a). This feature is at odd with what observed for major elements and leads to hypothesize that the Fe, K and Ca differences between the two groups are not necessarily primary features, but have been possibly acquired during post-magmatic alteration of the rock. Weathering of feldspars, amphibole and other mafic phases at different degrees may result in inter-element fractionation of these three elements, resulting in the apparently distinct liquid lines of descent observed in Figure 8.

As a whole, andesitic dykes are too much enriched in Mg#, for a given SiO₂ content, to be interpreted as partial melts of crustal rocks. Their high Mg# (0.64-0.80) is likely the result of the replacement of clinopyroxene by high-MgO chlorite. The Serre andesites show features related to the interaction between mantle-derived magmas which have variably interacted with crustal components, as reflected by variable LREE/HFSE (e.g., La/Nb = 2-13) and LILE/HFSE (e.g., Ba/Nb = 9-200) ratios. These variations might be caused by either magma mixing, crustal contamination as well as by concurrent assimilation and fractional crystallization (AFC; De Paolo, 1981). What now must be established is to understand if the interaction with crustal rocks (typically characterized by low HFSE, high LILE, high LILE/HFSE and high Pb; e.g., Rudnick and Gao, 2004, and references therein) did happen at mantle depths (e.g., after recycling of crustal rocks along subduction zones) or at crustal depths, during interaction of mantle melts with crustal lithologies (e.g., in magma chambers, via AFC processes).

To better understand the mechanisms that originated the two groups of intermediate Serre dykes, FC, AFC and mixing calculations have been carried out, by using the FC-AFC-FCA and mixing modeller program of Ersoy and Helvaci (2010). Results confirm that a single evolutionary trend for both andesite groups is very unlike. Indeed, in the Sr/Y vs. Y diagram (Figure 13a) calculations performed with A1a sample chosen as starting composition, indicate that the A andesites are genetically related each other by fractional crystallization (black solid line) of $\sim 20\%$ clinopyroxene, $\sim 20\%$ amphibole and $\sim 60\%$ plagioclase, starting from the average modal proportion of the most primitive sample. On the contrary, most of the F samples plot on the trajectory depicting mixing processes (black dotted line) between A1a-like parent magma and melts deriving from melting of Calabrian lower crustal metapelites (LCM; Del Moro et al., 2000; Figure 13a), now exposed in the northern sector of the Serre Massif. However, other samples of the F group do not plot along well defined trends, possibly reflecting variable interplay of mixing and FC-AFC processes, as well as possible minor alteration effects. The same calculations were replicated, considering F andesitic rocks as a separate population and F3 sample as the starting composition (Figure 13a). The modeled mixing curve (grey dotted line) nearly retraces the previous mixing trend, but in this case, simple fractionation (FC; grey solid line) from the more primitive F andesites appears to be the process more appropriate to explain compositions of the samples plotting out of the mixing trend. In addition, in the Ba/Nb vs. La/Nb diagram (Figure 13b), F dykes clearly separate from A ones, showing an AFC-mixing trend that seems to exclude any genetic link between the two dyke populations and highlights, instead, a significant contribution of crustal components, either as rockcontaminant and/or crustal melt in the generation of the F dykes. Alternatively, the Ba/Nb vs. La/Nb plot can be interpreted in a different way: elements resistant to post-magmatic hydrothermal alteration (e.g., REE and HFSE) are similar in the two groups (F and A dykes have overlapping La/Nb), with the most important differences seen only when elements mobile in fluid phases (e.g., LILE) are compared to relatively immobile elements (HFSE). According to this view, the different Ba/Nb could be also interpreted as the secondary effect of weathering, more evident in the A dykes than in the F samples. The overlapping Cr and Ni content of the two andesitic rock groups can be



Figure 13. a) Sr/Y vs. Y and b) Ba/Nb vs. La/Nb diagrams for Serre andesitic rocks. Black lines refer to A samples, grey lines refer to F samples. Serre lower crust metapelites (LCM; average composition) are from Del Moro et al., 2000. A1a - F3: starting composition. r = assimilation/fractionation ratio.

considered as a proof that substantial assimilation did not occur, because this would have caused cooling of the hybrid magma with the following increase of fractional crystallization of Ni-Cr-rich mafic phases (e.g., olivine and pyroxenes).

Nevertheless, the high Ba/Nb (25-201), La/Nb (3.1-6.2) and Th/La (0.25-0.42) and the low Nb/La (0.16-0.32) ratios of the A group dykes are a clear evidence of involvement of crustal lithologies in the genesis of these samples. However, the straightforward results of the geochemical modeling indicate that simple fractional crystallization from a A1a-like parent magma may be considered the main petrogenetic process for these andesites.

FC-AFC calculations have been carried out also for the dacite-rhyodacite dykes. Results (not shown) suggest the absence of fractional crystallization links with the andesitic melts. The geochemical similarities between the felsic dykes and the Serre granitoids can be interpreted hypothesizing that the high-K felsic volcanic rocks are related to partial melting of crustal rocks (e.g., von Blanckenburg et al., 1998; Altherr et al., 2000; Altherr and Siebel, 2002). At this purpose, crustal involvement in the origin of dacitic and rhyodacitic compositions is investigated using the results of dehydration melting experiments of different crustal rocks, such as amphibolites, metagreywackes tonalitic gneisses. and metapelites, under variable melting conditions (Altherr and Siebel, 2002, and references therein). Figures 14a-b, ST dacitic-rhyodacitic In compositions are broadly compatible with an origin by melting of a metapelitic source. Nevertheless, these diagrams highlight the already reported separation in the two sub-groups of silica-poor and silica-rich ST samples, suggesting a derivation from different crustal sources. A metapelitic crustal source appears to be likely for the silica-rich magmas, while a two-components metapelitic-metabasaltic sources seems to have been involved in the genesis of the silica-poor ST dykes.

Moreover, in ST silica-poor samples, Na₂O contents and Mg# values are respectively higher and lower than those from silica-rich samples, consistent with the assumption that partial melts from metabasaltic sources are generally characterized by higher contents of Na₂O and lower values of Mg# than melts from other sources (Rapp and Watson, 1995, and references therein). Additionally, the distinct petrographic features observed in ST silica-rich samples (e.g., presence of magmatic white mica), also support a different genetic mechanism.

The composition of group VZ dacitic dykes appears consistent with the derivation from a metapelitic crustal source in the Mg# vs. SiO₂ diagram (Figure 14a), whereas in the Al₂O₃/(MgO + FeO) vs. CaO/(MgO+FeO) diagram (Figure 14b), VZ samples show a very scattered distribution that can be hardly related to a pure metapelitic crustal source composition. Nevertheless, even considering the alteration effects, the composition of the dacites could results from hybridization of mantle derived melts with metapelites. Such mixing processes between crustal- and mantle-derived melts appear consistent with the "disequilibrium" textures observed in the VZ dacitic rocks (mantled feldspar phenocrysts, quartz ocelli and dissolution textures).

Conclusions

Post-collisional magmatic dykes, intruding metamorphic basement rocks as well as late Hercynian granitoids in four different areas of the Serre Massif, southern Calabria-Peloritani Orogen, range in composition from andesites (A-F groups) to dacites-rhyodacites (VZ-ST groups), all showing a medium- to high-K calcalkaline affinity. The two groups of andesites show subtle major element differences (mostly Fe, Ca, and, in a lesser amount, Al and K) that can be alternatively interpreted as different parental magma compositions or, more likely, as



Figure 14. Composition of Serre dacite-rhyodacites compared with composition of partial melts obtained in experimental studies by melting of various crustal rocks. a) SiO_2 vs. Mg# diagram. b) molar CaO/(MgO + FeO) vs. molar $Al_2O_3/(MgO + FeO)$ diagram. (Altherr and Siebel, 2002, and references therein).

secondary effects due to the diffuse postmagmatic weathering. Relatively immobile incompatible and compatible trace element content and ratios of the two andesite dyke groups are very similar, the only differences being found only for relatively mobile incompatible trace elements (e.g., LILE). These features can be interpreted as evidence of crustal contamination at shallow depths starting from a similar source or, more likely, as post-magmatic elemental mobility. Two sub-groups of silica-rich and silica-poor rocks, have been detected among the ST daciterhyodacite samples, that define separate behaviours in variation trends, reflecting further different petrogenetic mechanisms.

All the intermediate-felsic Serre dykes show subduction-related geochemical typical signatures, as typically found in other postcollisional igneous rocks of the European Hercynian Belt. On the whole, the studied calcalkaline dykes, intruding Hercynian phyllites and late-Hercynian undeformed granitoid rocks, were likely generated in a post-collisional extensional context, characterized by lithosphere upwelling thinning promoting of hot asthenosphere. This is indeed the general framework envisaged for dykes of roughly similar composition widespread in western Europe, such as those from the Sardinia-Corsica Domain (Atzori and Traversa, 1986; Atzori et al., 2000; Traversa et al., 2003), for which a transition from a compressive to extensional geodynamic setting has been invoked to explain their geochemical features. Calcalkaline affinity and geochemical signatures of such postcollisional rocks have been interpreted as a result of variable crustal contamination of mantle magmas derived from subduction-modified lithospheric and/or asthenospheric mantle sources.

Geochemical compositions of Serre andesitic dykes (A and F groups), including relatively high Mg# values and MgO, Ni and Cr contents, as well as variable LREE/HFSE and LILE/HFSE ratios and LILE and LREE enrichment, are strongly indicative of both mantle and crustal contributions. Results of the FC-AFC calculations suggest that interaction with crustal component took probably place at mantle depths, in the case of the group A andesites, through mantle metasomatism by subduction-related fluid/melts of crustal derivation. Partial melting of this enriched mantle source produced the most primitive group A melts that later evolved by simple fractional crystallization processes. Group F andesites were instead produced by AFC processes taking place during stalling and fractionation of mantle-derived melts in the crust and concurrent assimilation of wall rocks similar to the lower crustal Serre metapelites.

Group VZ dacites and silica-poor ST rocks likely resulted from hybridization, at variable extent, of basaltic mantle magma with pelitic metasediment. Finally, a purely crustal origin by partial melting of a metapelitic source is envisaged for the most acidic dykes, namely the ST silica-rich rhyodacites. Geochemical withingroup variations are in this case best modelled in terms of variable entrainment of restite components in the produced melts.

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