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Gabbroic xenoliths in tuff-breccia pipes from the Hyblean Plateau: insights into the nature and composition of the lower crust underneath South-eastern Sicily, Italy

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Summary

Crust-derived xenoliths hosted by Miocene basaltic diatremes in the Hyblean Plateau (south-eastern Sicily, Italy) provide new information regarding the nature of a portion of the central Mediterranean lower crust. These xenoliths can be divided into three groups: gabbros (plagioclase + clinopyroxene + Fe–Ti oxides \pm apatite \pm amphibole \pm Fe-rich green spinel), diorites (An-poor plagioclase, clinopyroxene \pm Fe–Ti oxides \pm orthopyroxene) and mafic granulites (plagioclase + clinopyroxene + green spinel \pm orthopyroxene \pm Fe–Ti oxides). Gabbros form the main subject of this paper. They represent cumulates whose igneous texture has been locally obliterated by metamorphic recrystallization and shearing. They were permeated by Fe–Ti-rich melts related to tholeiitic-type fractional crystallisation. Incompatible element ratios (Zr/Nb = 5–26; Y/Nb = 1.4–11) indicate that these cumulate gabbros derived from MORB liquids. Late-stage and hydrothermal fluids caused diverse, sometimes important, metasomatic trasformations. Petrographic and geochemical comparison with gabbroids from well-known geodynamic settings show that the Hyblean lower crustal xenoliths were probably formed in an oceanic or oceanic-continent transition environment.

Introduction

To define the mineralogical and geochemical characteristics and the structure of the lower continental crust, petrologists have studied granulite terrains outcropping on the Earth's surface and lower crustal xenoliths brought to the



Fig. 1. A Main structural domains of Sicily and neighbouring areas (simplified after *Lentini* et al., 1996); **B** Geological sketch map of the Hyblean area and location of the main xenolith occurrences

surface by ascending basaltic and kimberlitic magmas (*Downes*, 1993; *Rudnick*, 1992).

A large number of lower crustal xenoliths have been recently found in a few diatreme pipes in the Hyblean Plateau, in the south-eastern corner of Sicily (Southern Italy; Fig. 1). The Hyblean lower crust has been traditionally considered to belong to the African continental crust (*Burollet* et al., 1978: *Lentini* et al., 1996; *Catalano* et al., 2000). This conclusion is largely based on the interpretation of seismic data. However, only one seismic line crosses the western margin of the Hyblean Plateau (*Carrozzo* et al., 1987) and a few other refraction and reflection profiles transect only neighbouring areas (*Catalano* et al., 2000). Moreover, the interpretation of seismic soundings is closely related to a chosen lithological model. For instance, experimental results show that partially serpentinized peridotites have seismic wave velocities (V_p , V_s) similar to granitoids and other felsic rocks (e.g. *Hyndman* and *Peacock*, 2003).

Positive Bouguer anomaly and strong magnetic anomalies (up to 4500 cgs) have been interpreted as due to the presence of large mafic igneous masses just below the sedimentary-volcanic Meso-Cenozoic succession (*Arisi Rota* and *Fichera*, 1987; *Ciminale* and *Wasowski*, 1989). According to *Vai* (1994), the distribution of pelagic deposits within the southern Mediterranean area suggests that the Hyblean Plateau may represent the western part of the Permo-Triassic Palaeo-Tethys ocean. Moreover, *Sapienza* and *Scribano* (2000) underlined the absence among the xenoliths of lithologies typical of the continental middle-upper crust (e.g. granitoids, felsic granulites, schists) and the overall basaltic composition of the xenoliths. The choice of the one hypothesis (continental crust) over the other (oceanic crust) has important geological implications, i.e. on the location of the boundary between the African and European plates, on the nature of the eastern border of the Pelagian block (Fig. 1) and, more generally, on the geodynamic evolution of the Central Mediterranean area.

In this paper we present new whole rock (major and trace elements) and mineral chemistry (major elements) data of a suite of lower crustal xenoliths collected from the Hyblean Plateau, giving major emphasis to the gabbroic lithotypes. We also incorporate published data (*Scribano*, 1988; *Mazzoleni* and *Scribano*, 1994; *Tonarini* et al., 1996; *Punturo*, 1999; *Sapienza* and *Scribano*, 2000; see Table 1). The main aim of this study is to understand the origin and original tectonic setting of the deep Hyblean crust. It is opportune to refer to the case of Pannonian Basin (*Embey-Isztin* et al., 2003) as an example of a study that started with the assumption that the xenoliths were continental in origin, having instead produced the surprising result that that region is underlain by rocks of oceanic derivation.

Geological setting

The island of Sicily is located in the central Mediterranean area, in a geodynamic environment related to the collisional movement between the European and African Plates (*Dewey* et al., 1989). In the central Mediterranean area, *Lentini* et al. (1996) described three main structural domains (Fig. 1): (a) the *hinterland domain*, i.e. Tyrrhenian Basin and Sardinia-Corsica block; (b) the *orogenic domain* consisting of the Apenninic-Maghrebian thrust system and (c) the *foreland domain*, which represents undeformed crustal sectors not involved in the Apenninic-Maghrebian orogeny, i.e. Pelagian block and Ionian basin. The latter is generally considered to be an undeformed relict of oceanic crust of probable Jurassic age (e.g. *Finetti*, 1982). Recently *Catalano* et al. (2000), interpreting seismic reflection profiles, have distinguished two Ionian crustal sectors: the western, showing transitional-type crust and the eastern, consisting of old oceanic lithosphere.

The Hyblean Plateau is an emergent portion of the Pelagian block. It is in contact with the Gela nappe, belonging to the Apenninic-Maghrebian thrust system (north-western) and the Malta Escarpment (east) which separates the Plateau from the Ionian Basin. The exposed part of the Hyblean sedimentary sequence consists of Upper Cretaceous to Miocene deep-water carbonates and Plio-Quaternary open-shelf clastics, with intercalation of different levels of basic volcanic rocks (*Bianchi* et al., 1987; Fig. 1). Commercial boreholes have reached Triassic carbonates and basic igneous rocks at about 3 km b.s.l., and geophysical data suggest that this entire sequence is 8–10 km thick (*Chironi* et al., 2000). Two main fault systems cross-cut the Hyblean area. One, trending NE–SW, is extensional; the other, trending NNW–SSE, mostly consists of strike-slip faults (*Grasso* and *Reuther*, 1988). Three main magmatic episodes have been recognised: Cretaceous alkali-basalts (*Romano* and *Villari*, 1973), upper Miocene alkali-basalts and tuff-breccia pipes (*Cristofolini* et al., 1981; *Carbone* and *Lentini*, 1981) and Plio-Pleistocene lavas. The Plio-Pleistocene lavas are tholeiitic to alkaline basalts, basanites and rare

nephelinites. They are primary liquids whose different compositions depend on the degree of partial melting, the different depths and the metasomatic enrichments of the source (*Beccaluva* et al., 1998). Interpretation of the Sr–Nd–Pb isotopic data for the Upper Miocene and Plio-Pleistocene lavas has excluded the presence of continental crust contamination (*Trua* et al., 1998; *Bianchini* et al., 1999).

Petrography

Ultramafic and feldspar-bearing xenoliths occur in Miocene pipes of the Hyblean Plateau. The tuff-breccia deposit from Valle Guffari (*vg*; Fig. 1) is the most significant xenolith occurrence in the region. The ultramafic suite consists of mantle-derived material, represented by spinel lherzolites, spinel harzburgites and various pyroxenite types, i.e. Cr-diopside websterites, Al-augite websterites, spinel and garnet clinopyroxenites (*Scribano*, 1987; *Nimis* and *Vannucci*, 1995; *Punturo* et al., 2000).

The feldspar-bearing suite, which is considered to be derived from the Hyblean lower crust, consists of sub-rounded xenoliths 5 to 15 cm in diameter. They lack evident infiltrations of the host basaltic melt. On the basis of texture and mineral association, we have subdivided them into three main groups, whose relative proportions are roughly estimated from the number of collected xenoliths: *i*) mafic granulites, $\approx 60\%$, *ii*) gabbros, $\approx 30\%$, and *iii*) diorites, $\approx 10\%$. Both gabbros and diorites show different degrees of shearing, recrystallization and metasomatic transformations. The gabbros are the subject of this study.

Gabbros

Plagioclase, clinopyroxene, Fe–Ti oxides, green hercynitic spinel, apatite and rare red-brown amphibole form the gabbro paragenesis. These still preserve their igneous texture in spite of having undergone shearing and more or less pronounced metamorphic re-equilibration.

Ductile and brittle deformation textures are widespread in these xenoliths. The plagioclase porphyroclasts (Pl I) show either crystal-plastic deformation or notable grain-size reduction due to recrystallisation (plagioclase neoblasts: Pl II; Fig. 2a,b). Conversely, most of the pyroxenes show brittle deformation (clinopyroxene

Fig. 2. Textural features of xenoliths from Hyblean diatremes. **a** relic plagioclase porphyroclast (Pl I) within the matrix formed by re-crystallised clinopyroxene (Cpx II) and plagioclase (Pl II) in oxide-rich gabbros (crossed nicols); **b** grain-size reduction along shear plane between porphyroclasts in oxide-rich gabbros (plane light); **c** anhedral Fe–Ti oxides III related to glassy pockets within the oxide-rich gabbro matrix; deformed sub-rounded Cpx I is visible (plane light); **d** sheared texture in oxide-rich gabbro (plane light). Plagioclase appears white, clinopyroxene grey, Fe–Ti-rich oxide (III) black; **e** corrosion edge in plagioclase porphyroclasts (Pl I) in contact with glassy pockets (plane light); **f** scanned polished surface of a sheared diorite xenolith FB-h3; **g** polished surface of diorite xenolith FB-h2 showing modal layering and brittle fractures. Mineral whitish to pale grey is plagioclase, medium-grey is pyroxene. Full explanation is given in the text; **h** granoblastic texture in mafic granulite (plane light)

porphyroclasts: Cpx I) rarely surrounded by pyroxene neoblasts (Cpx II) (Fig. 2b, c, d). Cpx I shows abundant ilmenite exsolution lamellae, while Cpx II is exsolution-free (Fig. 2c).

Fe-Ti oxides occur included in silicates (Fe-Ti oxides I; Fig. 2b), as scattered grains (2–5 vol%; Fe-Ti oxides II) interstitial to silicates or, most commonly, as anhedral amoeboid-shaped grains, forming large opaque patches (8–18 vol%;



Fe–Ti oxides III) within the grain frame of the gabbro (Fig. 2c, d, e). Locally, anhedral micrograins of green hercynitic spinel are found within these patches. On the basis of the Fe–Ti oxide modal abundances, the gabbros can be subdivided into two subgroups: the oxide-rich gabbros (Fe–Ti oxides $\geq 8 \text{ vol}\%$) and oxide-poor gabbros (Fe–Ti oxides $\leq 8 \text{ vol}\%$).

The Fe–Ti oxides III are generally associated with melt pockets. These pockets of altered, turbid silicate glass with fine-grained clinopyroxene, plagioclase and opaque ores are ubiquitous and especially abundant in the oxide-rich gabbros. They show disequilibrium relationships with the neighbouring silicates: Pl I grains in contact with the glass are strongly corroded (Fig. 2e), while Cpx I grains exhibit a thin purplish rim (Fig. 2c). Zeolites fill vugs occuring in these pockets. The volume of melt pockets, and thus Fe–Ti oxide III, is positively correlated with the degree of deformation of the host rocks. A number of secondary minerals (i.e. Fe–Mg amphiboles, chlorites, smectites, Na-rich zeolites, calcite and Fe-hydroxides; Fig. 2c, e) fill both vugs and veinlets or locally replace primary minerals.

Diorites

Diorites are coarse- to medium-grained leucocratic rocks. Their paragenesis is dominated by Na-rich plagioclase with subordinate clinopyroxene and Fe–Ti oxides. They show variable microstructures.

The most common type shows an evident lineation (Fig. 2f) due to the alignment of finely granulated pyroxene relics in a matrix of medium- to fine-grained feldspar crystals. Locally, a double shell of Fe–Ti hydroxides (outer shell) and calcite (inner shell) separates the pyroxene grains from the feldspar matrix. In some cases aggregates of polycrystalline calcite replace the pyroxene. A slightly pleochroic bronzite occurs in these rocks, up to 6 vol%, either as anhedral isolated grains or tiny, corroded relics surrounded by calcite.

Another type of diorite, coarser grained than the first variety and partially recrystallized, shows modal layering due to alternating 3–5 cm thick layers rich in feldspar and thinner, darker layers rich in pyroxene and opaques. Pyroxene grains are often rimmed and locally replaced by a yellowish turbid material consisting of fine aggregates of secondary minerals hosting tiny vugs filled with analcite and zeolites. The same material also fills brittle micro-fractures occurring in the rock as subparallel sets discordant to the layering (Fig. 2g).

Plagioclase from the diorite xenoliths is affected by different degrees of alkaline metasomatism in the form of albite and (subordinate) K-feldspar veins (*Scribano* et al., 2003). In addition, chlorites, zeolites, smectites, Fe-oxide/hydroxides and carbonates are ubiquitous in these rocks, partially replacing the igneous minerals.

In a few cases, the metasomatic transformations and deformations have totally obliterated the original mineral compositions and textures. Fractured relics of alkali feldspar ($An_3-Ab_{80}-Or_{17}$), surrounded by tiny rounded or elongated feldspar neoblasts, are embedded in an abundant matrix of chlorites and smectite. Rare irregular hematite grains and a number of tiny opaque patches also occur in the phyllosilicate matrix. The alkali feldspar porphyroclasts exhibit chessboard extinction, 82–85% triclinicity and a low-temperature structural state (*Scribano* et al., 2003). In one of these metasomatic rocks (sample FBZ) several monazite

and zircon grains have been found in the chlorite patches. These metasomatic rocks closely resemble the hydrothermal zircon-bearing, alkali feldspar – chlorite "blackwalls", enveloping rodingite bodies from ophiolite massifs (e.g. Sudetic ophiolite, SW Poland: *Dubinska* et al., 2004). A detailed mineralogic and geochemical study on these metasomatic rocks will be reported elsewhere, though the preliminary results supplied by *Scribano* et al. (2003) will be taken into account for the present discussion.

Mafic granulites

Scribano (1988) gave a preliminary report on the petrography of the mafic granulites. The following summarises their principal features. Hyblean granulites show granoblastic microstructures with fine (0.3-1 mm) to medium (0.5-3 mm) grain sizes (Fig. 2h). They consist of plagioclase, clinopyroxene, orthopyroxene and green Al-spinel. Generally, clusters of pyroxenes and plagioclase grains are randomly distributed in the rocks. Rarely, mafic and felsic minerals are arranged in alternating irregular layers visible at the hand-specimen scale. The occurrence of rare spinel-bearing anorthositic granulites and even rarer feldspar-bearing pyroxenites have also been reported. Fine exsolution lamellae of Ca-poor from Ca-rich pyroxene and vice-versa are common in the pyroxenes. Spinel occurs as wormshaped intergrowths in the pyroxene grains, as discrete grains between the coexisting silicates and more rarely as "droplets" enclosed in the feldspars. Striking coronitic textures and trails of CO₂ fluid inclusions are common in these rocks (*Scribano*, 1988; *Pompilio* and *Scribano*, 1992).

Analytical techniques

Table 1 reports the sample set considered. To supplement the existing data on the mafic granulites, representative samples of oxide-rich and oxide-poor gabbros and diorites were selected. Chemical analyses of minerals were performed at the C.N.R. Instituto di Geoscienze e Georisore (I.G.G.), Padova, using a Cameca Camebax equipped with four wavelength-dispersive spectrometers. Operating conditions were 15 kV and 15 nA. Synthetic oxides and minerals were used as standards. X-ray intensities were automatically corrected to oxide percent concentrations by on-line PAP procedure. Additional analyses were performed using a Cambridge Instrument SEM fitted with an EDS Link (operating conditions: 1.5 kV, 20 nA, lifetime 100 s, ZAF corrections) at the Istituto Nazionale di Geofisica e Vulcanologia – Catania.

Geochemical analyses were performed at Service d'Analyse des Roches et des Minéraux (SARM) laboratories in Nancy, France. Whole-rock major elements were obtained by Inductively Coupled Plasma Emission Spectrometry (ICP), trace elements by Inductively Coupled Plasma Mass Spectrometry (ICP-MS). CO_2 was detected volumetrically. Total iron is reported as Fe_2O_3 . Detailed information on analytical methods and detection limits are reported in: www.crpg.cnrs.nancy.fr/SARM.

The whole rock δ ¹⁸O values have been obtained at Activation Laboratories (Ancaster, Ontario). Oxygen was extracted from powders of whole rocks by

Table 1. Summary of Hyblean lower crustal samples included in this study, references and provenance sites. Vg Valle Guffari; Cm Cozzo Molino (see Fig. 1). cpx clinopyroxene; pl plagioclase; ox Fe–Ti oxides; spl Al-spinel

Sample	Data	Ref*	Area
Ia) Oxide-r	ich gabbros		
XLP1	Major, trace, REE elements, cpx, ox and pl compositions	1	Vg
FB-f4	Major, trace, REE elements, cpx, ox and pl compositions	1, 2	Vg
FB80	Major, trace, REE elements, cpx, ox and pl compositions	1	Vg
FB-f31	Major, trace, REE elements, cpx, ox and pl compositions	1, 2	Vg
FB50	Major, trace, REE elements, cpx, ox and pl compositions	1	Vg
VB5	Major, trace, REE elements, cpx, ox and pl compositions	1	Vg
ML17	Major, trace, REE elements	3	Vg
FB-f1	Major, trace, REE elements, cpx, ox and pl compositions	1, 2	Vg
Ib) Oxide-p	poor gabbros		
FB70	Major, trace, REE elements, cpx, ox and pl compositions	1	Vg
FB-f7	Major, trace, REE elements, cpx, ox and pl compositions	1, 2	Vg
FB13	Major, trace, REE elements, cpx, ox and pl compositions	1	Vg
FBf32	Major, trace, REE elements	1	Vg
FB11	Major, trace, REE elements, cpx, ox and pl compositions	1	Vg
FBe11	Major, trace, REE elements	1	Vg
II) Diorites			
FB12	Major, trace, REE elements, cpx, ox and pl compositions	1	Vg
FB8	Major, trace, REE elements, cpx, ox and pl compositions	1	Vg
FB-h2	Major, trace, REE elements, cpx, ox and pl compositions	1, 2	Vg
FB-h3	Major, trace, REE elements, cpx, ox and pl compositions	1, 2	Vg
FBY	Major, trace, REE elements	1	Vg
III) Mafic g	granulites		
MVG1	Major, trace, REE elements; cpx, spl and pl compositions	4, 5	Cm
MVG2	Major, trace, REE elements; cpx, spl and pl compositions	4, 5	Cm
MG3	Major, trace, REE elements; cpx, spl and pl compositions	4, 5	Cm
MG4	Major, trace, REE elements; cpx, spl and pl compositions	5, 6	Cm
MG7	Major, trace, REE elements; cpx, spl and pl compositions	5, 6	Cm

* 1: this study; 2: *Sapienza* and *Scribano* (2000); 3: *Punturo* (1999); 4: *Tonarini* et al. (1996); 5: *Scribano* (1988); 6: *Mazzoleni* and *Scribano* (1994)

reaction with BrF₅, converted to CO₂ in a graphite furnace and analysed by mass spectrometer. The isotopic composition of a sample is given as: $\delta_{\text{sample}} = (R_{\text{sample}}/R_{\text{standard}} - 1)^*$ 1000 in per milunit, where R is ¹⁸O/¹⁶O and the standard is SMOW.

Mineral compositions

Pyroxenes

Compositions of representative clinopyroxenes are reported in Table 2. Clinopyroxenes from the studied xenoliths are mostly diopsides and only those from sample

	Oxide-ric	h gabbros	Oxide-po	oor gabbros		Diorite			
Sample	FB80 P	FB50 P	FB11 N	FB13 N	FB70 N	FB8 N	FB12 N	FB3 N	FB3 N
Wt%									
SiO ₂	51.86	50.65	47.49	50.85	50.40	51.25	52.17	52.53	53.60
TiO_2	0.47	0.78	1.87	0.58	0.98	0.49	0.46	0.27	0.00
$Al_2 \tilde{O}_3$	3.47	4.27	7.80	4.07	5.04	3.30	2.57	2.11	0.71
Cr_2O_3	0.03	0.00	0.12	0.00	0.02	0.02	0.03	0.00	0.00
FeO	7.03	7.46	6.82	8.05	8.13	8.72	10.05	9.64	23.0.9
MnO	0.32	0.26	0.19	0.23	0.27	0.53	0.75	0.62	1.15
MgO	13.92	13.61	11.95	13.36	13.18	12.55	13.42	12.61	20.12
CaO	22.79	23.07	22.86	22.98	20.86	21.96	19.24	20.75	0.91
Na ₂ O	0.46	0.68	0.97	0.67	1.05	1.08	1.27	1.12	0.00
Total	100.35	100.78	100.05	100.79	99.93	99.90	99.96	99.65	99.58
Mg#	78	76	75	77	74	72	70	70	60

Table 2. Chemical composition of pyroxenes of Hyblean xenoliths. $Mg\#=Mg/(Mg+Fe^{2+}+Fe^{3+})^*100$

FB12 (diorite) are augites. Overall, their Mg# value ranges from 70 to 85. Clinopyroxenes from the mafic granulites show the widest ranges in Mg# (71–85), Al_2O_3 (3.5–9 wt%), TiO₂ (0.5–1.5 wt%) and Na₂O (0.7–1.5 wt%). No intra-granular variations were found in these clinopyroxenes.

Clinopyroxenes from the oxide-rich gabbros show a relatively narrow range of Mg# (73–78; Fig. 3). The cores of the Cpx I generally display $Al_2O_3 \sim 4 \text{ wt\%}$, TiO₂ $\sim 0.7 \text{ wt\%}$ and Na₂O $\sim 0.6 \text{ wt\%}$, while their outermost rims in contact with the



Fig. 3. An (mol%) of plagioclase *vs* Mg# of coexisting clinopyroxenes from the Hyblean gabbros and diorites. The symbols correspond to the average An of plagioclases and Mg# of coexisting clinopyroxenes per sample. The bars indicate the variations within each sample

glassy patches have higher Al₂O₃ and TiO₂ contents. The diorite clinopyroxenes have the lowest average Mg# (69–71; Fig. 3) and show some variations in Al₂O₃ (2–4 wt%), TiO₂ (0.4–0.7 wt%) and Na₂O (1–1.5 wt%). The average Mg# of clinopyroxenes (Cpx II) from oxide-poor gabbros (Mg# = 74–75; Fig. 3) is intermediate between those of oxide-rich gabbros and diorite clinopyroxenes and partially overlaps the former group. Scarce Ca-poor pyroxene (Mg# = 60; CaO \approx 90 wt%), coexisting with a calcic one (Mg# = 70; CaO \approx 21 wt%), has been found only in recrystallized diorites (e.g. FB-h3: Table 1). Orthopyroxene from mafic granulites shows Mg# = 65–68, 2.0–6.5 Al₂O₃ wt% and 0.2–0.7 CaO wt% (*Scribano*, 1988).

Feldspars

Feldspar compositions in gabbros and diorites are reported in Table 3. Generally, they are plagioclase varying from An_{55} to An_{80} in gabbro xenoliths (except in FB70 which is An_{40} ; Fig. 3), from An_{15} to An_{30} in diorites, and from An_{55} to An_{82} in mafic granulites. Plagioclase compositions are something homogeneous within each sample (Fig. 3). The Or content is negligible in gabbro and mafic granulite feldspars, whereas it varies from 2 to 10 mol% in those from diorites.

Oxides

The chemical compositions of representative oxides are reported in Table 4. Opaques included in clinopyroxene and plagioclase porphyroclasts (type I) are Ti-magnetite and ilmenite, while the interstitial oxides (type II) are ilmenite.

	Oxide-ri	ch gabbro	Oxide-p	poor gabbre	0			Diorite	
Sample	FB80 P core	FB50 P rim	FB11 N	FB70 P core	FB70 P rim	FB13 P core	FB13 P rim	FB12 N	FB8 N
Wt%									
SiO ₂ TiO ₂	50.11 0.02	51.08 0.00	49.85 0.01	57.99 0.03	58.32 0.02	47.34 0.02	49.48 0.01	64.64 0.02	60.04 0.01
Al_2O_3 Cr_2O_3	31.50 0.05	30.93 0.00	31.51 0.00	26.03 0.00	26.41 0.02	32.68 0.00	32.13 0.07	21.85 0.00	24.43 0.00
FeO MnO	$0.06 \\ 0.01$	0.13 0.00	0.02 0.00	0.03 0.00	$\begin{array}{c} 0.05 \\ 0.01 \end{array}$	$0.17 \\ 0.00$	$\begin{array}{c} 0.08 \\ 0.06 \end{array}$	$\begin{array}{c} 0.04 \\ 0.00 \end{array}$	0.07 0.00
MgO CaO	0.00	0.01	0.01	0.01	0.00	0.00	0.00	0.00	0.02
Na ₂ O	3.19	3.81	3.14	6.65 0.29	6.79 0.27	2.12	2.95	9.06 1.56	7.66
BaO	0.000	0.000	0.05	0.29	0.027	0.02	0.16	0.00	0.47
Total	99.91	100.20	99.75	99.41	100.48	99.42	100.84	100.34	99.39
Ab mol% An mol% Or mol%	27.9 71.9 0.2	32.8 66.9 0.3	27.1 72.6 0.3	58.3 40.1 1.6	58.2 40.3 1.5	18 81.9 0.1	25.3 74.3 0.4	76.8 14.6 8.6	66.3 31 2.7

Table 3. Chemical composition of plagioclases of Hyblean xenoliths. P porphyroclast; N neoblast

	Oxide-ri	ich gabbro				Oxide-p gabbro	oor	Diorite	
Sample Type	FB80 I	FB80 III	FB80 III	FB4 III	FB4 III	FB7 II	FB70 I	FB13 III	FB13 I
Wt%									
SiO ₂	0.08	0.03	0.03	0.08	0.06	0.06	0.00	0.00	0.00
TiO ₂	7.93	52.11	0.05	53.89	19.23	10.36	43.45	38.99	41.43
Al_2O_3	2.71	0.60	60.03	0.33	6.26	3.84	0.84	0.28	0.77
FeO	80.98	39.31	24.57	35.70	63.65	77.78	50.66	55.06	54.46
MnO	0.34	0.61	0.27	0.77	0.58	0.34	0.43	0.48	0.38
MgO	2.82	6.68	11.83	8.50	6.96	1.96	3.15	2.69	1.34
CaO	0.04	0.08	0.00	0.07	0.08	0.00	0.04	0.00	0.05
Cr ₂ O ₃	0.06	0.00	0.11	0.03	0.01	0.34	0.05	0.00	0.00
Total	94.96	99.42	96.89	99.37	96.83	94.68	98.63	97.50	98.43

Table 4. Chemical composition of Fe-Ti oxides of Hyblean xenoliths. Types (I, II, III) are explained in the text

The patch-forming oxides (type III), characteristic of, but not restricted to the oxide-rich gabbros, consist of a coarse intergrowth of ilmenite and Ti-magnetite and rare green hercynitic spinel (FeO $\sim 24 \text{ wt\%}$; Al₂O₃ $\sim 60 \text{ wt\%}$). Spinel from the mafic granulites displays Al₂O₃ content from 46 to 65 wt%, FeO from 15 to 39 wt%, MgO from 10 to 17.5 wt%. Cr₂O₃ content never exceeds 6.5 wt% (*Scribano*, 1988).

Amphiboles

Amphibole in the oxide-poor gabbro FB70 is a titanian pargasite (*Leake* et al., 1997) with Mg# = 63. TiO₂, Al₂O₃ and CaO contents are 3.7 wt%, 13.4 wt% and 11.7 wt%, respectively (Table 5). In the glassy pockets within the gabbro and diorite xenoliths,

Sample	FB70	FB70	
-	core	rim	
Wt%			
SiO ₂	40.45	40.56	
TiO_2	4.37	3.70	
Al_2O_3	13.34	13.40	
Cr_2O_3	0.00	0.01	
FeO	12.66	12.42	
MnO	0.17	0.29	
MgO	11.51	11.82	
CaO	11.40	11.69	
Na ₂ O	2.96	2.95	
K ₂ O	0.92	0.91	
Total	97.77	97.74	

Table 5. Chemical composition of amphibole of Hyblean oxide-poor gabbro FB70

SEM investigations showed the presence of actinolite $(Al_2O_3 = 0.9 \text{ wt\%}; CaO = 10 \text{ wt\%}; Mg\# = 60)$. Rare pargasite also occurs in the mafic granulites $(Mg\# = 80; TiO_2 = 2.7 \text{ wt\%}; Al_2O_3 = 16.5 \text{ wt.\%}, CaO = 11.4 \text{ wt\%}; Scribano, 1988).$

Whole-rock chemistry

New major and trace element data of Hyblean lower crustal xenoliths are reported in Table 6. The following discussion of the data refers to the entire sample set (Table 1). The whole rock composition of the host alkali-basalt (sample LOR) is also reported for comparison.

Major elements

Figure 4 shows the major element variations against MgO content of the Hyblean lower crustal xenoliths. The bulk composition of the gabbros is strongly mode-dependent: they show the widest variations in SiO₂ (from 36.4 to 49.6), Fe₂O₃ (from 11.5 to 22.7 wt%) and TiO₂ (from 2.9 to 5.5 wt%), driven by the modal abundance of Fe–Ti oxides.

The diorite xenoliths show the highest SiO₂ (up to 57 wt%) and alkali contents (Na₂O = 5.5–7.3 wt% and K₂O up to 2.5 wt%; always Na₂O > K₂O). This composition is related to the modal abundance of Na-plagioclase, which has been affected by pervasive alkaline metasomatism. Diorites also show low MgO and CaO contents. The gabbros and the mafic granulites have K₂O \leq 0.2 wt%, with the latter showing the lowest K₂O contents (0.04–0.09 wt.%). P₂O₅ is always <1 wt.% in all xenoliths, except for the apatite-rich oxide-rich gabbro ML17 (3.5 wt.%). The loss on ignition (LOI) in gabbros and diorites (1.9–6.2 wt.%) is higher than in mafic granulites (1.5–0.7 wt.%). CO₂ contents never exceed 0.5 wt%, except in some diorites whose clinopyroxene grains are partially replaced by calcite (e.g. 3.6 wt% in sample FB-h3).

Trace elements

Figure 5 shows chondrite-normalised Rare Earth Element (REE) patterns and N-MORB normalised spiderdiagrams of the Hyblean lower crustal xenoliths. The REE patterns of all gabbro xenoliths are flat from La to Sm or slightly LREE enriched (La_N/Sm_N ~0.7–1.7). They generally show a moderate positive Eu anomaly ($1 \le Eu/Eu^* \le 2$, where $Eu^* = (Sm_N + Gd_N)/2$; *Salters* and *Shimizu*, 1988). MREE are fractionated with respect to the HREE. Samples VB5 and Fbe11 show a similar pattern as the other gabbros except for the Eu anomaly. The pyroxene-rich and oxide-rich gabbro FB-f1 shows an upward-convex REE pattern with LREE depletion (La_N/Sm_N ~0.5).

The diorite xenoliths show a marked LREE/HREE fractionation $(La_N/Yb_N = 8.5-20.4)$ with a pronounced positive Eu anomaly (Eu/Eu^{*} up to 4.7), indicating that accumulation of plagioclase played an important role in the diorite formation. The mafic granulites show the lowest REE abundances, with LREE/HREE fractionation (La_N/Yb_N = 1.5-4.5). A positive Eu anomaly is also present.

In the N-MORB-normalised incompatible elements diagrams (Fig. 5), the gabbros and diorites are generally enriched from Rb to P, with evident Ba and Sr

ladie 0. /M	ajor ana	irace elem	teri comp	fo uomso	unanaku	wer crusic	и хепонных	. 11.a. <i>not</i>	anaiysea;	190 .L.D.O	now delection	inmu no:		
	Oxide-r	ich gabbro	S			Oxide-po	or gabbro			Diorites			Host lava	
	XLP1	FB50	FB80	VB5	FBf32	FB13	FB11	FBe11	FB70	FB8	FBΥ	FB12	LOR	
(wt%)														
SiO ₂	39.51	39.08	39.83	42.68	45.66	43.08	46.76	48.4	46.01	51.99	48.51	56.73	41.56	
TiO_2	4.73	4.03	3.8	2.89	2.23	2.41	0.96	1.26	2.19	1.43	2.49	1.52	2.29	
Al_2O_3	12.9	14.85	15.55	10.75	20.39	21.58	22.07	13.66	15.76	19.99	15.83	18.36	12.35	
Fe ₂ O _{3 tot}	16.97	15.22	14.90	12.75	6.48	9.9	4.57	9.93	10.01	5.40	9.27	5.01	10.78	
MnO	0.17	0.14	0.15	0.15	0.07	0.07	0.06	0.12	0.14	0.06	0.19	0.09	0.15	
MgO	6.73	7.43	7.28	9.77	5.3	4.27	5.71	9.14	6.64	3.61	3.61	2.67	11.61	
CaO	12.47	13.82	13.4	16.29	13.35	13.37	16.17	12.29	11.89	6.35	8.17	4.23	9.83	
Na_2O	2.23	1.59	1.23	1.66	2.5	2.37	2.03	2.44	3.01	5.8	5.48	6.03	1.85	
K_2O	0.13	0.22	0.16	0.16	0.17	0.12	0.09	0.18	0.23	0.6	1.16	2.46	1.15	
P_2O_5	0.36	0.17	0.08	0.34	0.18	0.25	0.12	0.06	0.34	0.11	0.76	0.35	1.15	
LOI	3.37	3.58	2.49	2.61	3.72	2.63	1.9	2.41	3.64	4.57	4.6	2.67	7.25	
H_2O	n.a.	3.01	2.35	2.36	n.a.	2.49	1.64	n.a.	3.16	4.17	1.28	2.53	n.a.	
CO_2	n.a.	0.57	0.14	0.25	n.a.	0.14	0.26	n.a.	0.48	0.4	3.32	0.14	n.a.	2
tot	99.57	100.13	98.87	100.05	100.05	100.05	100.44	99.89	99.86	99.91	100.07	100.12	99.97	
$Mg^{\#}$	48.02	53.22	53.23	64.1	65.58	50.14	74.44	68.18	60.70	60.89	47.57	55.37	70.75	
La (ppm)	8.34	6.03	6.79	6	5.04	7.21	3.7	4.08	17.12	7.54	38.93	20.66	81.45	
Ce	22.84	14.83	16.19	23.34	13.29	15.38	9.52	11.4	39.78	14.22	85.06	44.14	151.91	
Pr	3.52	2.22	2.48	3.63	1.86	2.08	1.55	1.73	5.33	1.7	11.31	5.89	16.41	
Nd	18.34	11.57	12.38	18.86	9.39	10.17	8.64	9.4	24.49	7.36	49.42	26.45	60.69	
Sm	4.84	3.02	3.49	5.26	2.4	2.37	2.39	3.26	5.32	1.8	10.25	5.7	11	
Eu	2.11	1.37	1.48	2.01	1.36	1.17	1.14	1.32	2.39	1.28	5.17	4.34	3.36	
Gd	4.75	3.15	3.48	5.31	2.33	2.25	2.46	3.92	4.91	1.62	8.47	4.86	9.77	
Tb	0.66	0.44	0.49	0.73	0.32	0.31	0.31	0.6	0.7	0.24	1.1	0.65	1.2	
Dy	3.56	2.32	2.66	4.07	1.74	1.65	1.72	3.51	3.79	1.33	5.59	3.44	6.22	
													(continued)	

Gabbroic xenoliths from the Hyblean Plateau

Host lava $\begin{array}{c} 1.06\\ 2.54\\ 0.33\\ 1.92\\ 1.92\\ 1.92\\ 0.29\\ 8.6\\ 8.6\\ 8.6\\ 8.6\\ 8.6\\ 5.3\\ 5.3\\ 5.3\\ 995\\ 5.3\\ 208\\ 228\\ 2353\\ 3353\\ 302\\ 302\end{array}$ LOR FB12 FBY Diorites $\begin{array}{c} 0.24\\ 0.59\\ 0.09\\ 0.49\\ 0.49\\ 0.06\\ 0.13\\ 0.38\\ 0.13\\ 0.38\\ 0.38\\ 0.13\\ 0.38\\ 0.13\\ 0.38\\ 0.13\\ 0.55\\ 0.13\\ 0.55\\ 0.13\\ 0.55\\ 0.13\\ 0.55\\ 0.13\\ 0.55\\ 0.13\\$ FB8 0.66 1.75 0.22 1.31 0.29 0.09 1.2 0.09 1.2 0.09 1.2 0.09 1.2 0.09 1.5 35 55 FB70 FBe11 **Dxide-poor** gabbro 0.29 0.68 0.09 0.07 0.07 3.2 0.07 0.07 0.07 1541 0.71 1541 0.71 1541 0.71 18 7.8 231 231 231 233 79 FB11 $\begin{array}{c} 0.27\\ 0.63\\ 0.08\\ 0.048\\ 0.07\\ 5.6\\ 0.1\\ 6.23\\ 0.1\\ 0.1\\ 0.37\\ 0.$ FB13 FBf32 $\begin{array}{c} 0.69 \\ 1.65 \\ 0.21 \\ 0.21 \\ 1.28 \\ 0.18 \\ 0.16 \\ 0.16 \\ 5 \\ 0.44 \\ 0.16 \\ 5 \\ 0.44 \\ 0.16 \\ 5 \\ 2.2 \\ 1.8 \\ 1.8 \\ 4.0 \\ 1.8 \\ 1.8 \\ 1.8 \\ 1.8 \\ 1.8 \\ 1.8 \\ 1.8 \\ 1.1 \\ 1.8 \\ 1.1 \\ 1.8 \\ 1.1 \\ 1.8 \\ 1.1 \\$ VB5 $\begin{array}{c} 0.48\\ 1.07\\ 0.14\\ 0.14\\ 0.14\\ 0.12\\ 0.12\\ 0.25\\$ FB80Oxide-rich gabbros $\begin{array}{c} 0.4 \\ 0.96 \\ 0.12 \\ 0.12 \\ 0.73 \\ 0.11 \\ 10 \\ 1.4 \\ 0.07 \\ 0.11 \\ 0.07 \\ 0.11 \\ 0.07 \\ 0.11 \\ 0.07 \\ 0.12 \\ 0.07 \\ 0.12 \\ 0.07 \\ 0.12 \\ 0.01 \\ 0.14 \\ 0.07 \\ 0.07 \\ 0.01 \\$ FB50 XLP1

V. Scribano et al.

76

 Table 6 (continued)



Fig. 4. Whole rock MgO *vs* major oxides for the Hyblean crustal xenolith. For comparison, the fields of gabbroids from different geodynamic settings are also shown

positive anomalies, and depleted in Zr–Hf and HREE. Oxide-rich gabbros patterns are characterised by positive Ti anomaly, in agreement with the relative high modal abundances of Fe–Ti-oxides. Mafic granulites show incompatible element patterns resembling those of the gabbros and diorites but a lower abundance and with a less marked Zr and Hf depletion.



Fig. 5. C1-normalized REE patterns (normalising values after *Anders* and *Grevesse*, 1989) and spiderdiagrams (N-MORB normalising values after *Hoffmann*, 1988; P after *Sun*, 1980) for the Hyblean crustal xenoliths. Open circle: host basalt

Oxygen isotopes

Two samples with no petrographic evidence of important alteration products of primary minerals were chosen for whole rock oxygen isotope analyses. The diorite FB12 has $\delta^{18}O_{SMOW} = 8.7\%$ (LOI = 2.67 wt.%, MgO = 2.67 wt%), and the oxide-poor gabbro FB11 has $\delta^{18}O_{SMOW} = 7.2\%$ (LOI = 1.9 wt%, MgO = 5.71 wt%).

Discussion

Origin of the different groups of Hyblean mafic xenoliths

The textural, mineralogical and geochemical heterogeneity observed in the Hyblean xenoliths suggests that the lower crust beneath south-eastern Sicily has experienced a complex petrological evolution. Petrographic data demonstrate the igneous origin of the gabbros. Their mode-controlled chemical compositions suggest that they represent cumulates. In fact, all studied rocks plot outside the primary basaltic liquids field in Mg# vs SiO₂/Al₂O₃ diagram (Fig. 6).

The term "cumulate" is here unrelated to any specific petrogenetic models, i.e. those invoking crystal settling in a large magmatic body (for reviews see *Cawthorn*, 1996). The evident relationship between Fe–Ti oxides III and melt



Fig. 6. (SiO_2/Al_2O_3) vs Mg# bivariate plots of Hyblean crustal xenoliths. Field of primary basaltic liquids and directions of mineral phase accumulation after *Kempton* et al. (1995). Same symbols as in Fig. 4

pockets (now glass or devitrified glass), the occurrence in primary phases of reaction rims and embayed microstructures against the melt pockets, and their clear connection with deformation textures, all imply the injection of strongly fractionated Fe–Ti-rich melts, probably filter-pressed from gabbroic or dioritic crystalmushes, within neighbouring gabbroic cumulate masses under shearing conditions. These melts reacted with the host mineral assemblage and precipitated the large oxide minerals. The Fenner-type fractionation trend leading to the oxide-related melts suggests a tholeiitic nature for their parent magma.

Such magmatic infiltration has been documented in lower crustal xenoliths of the Cima volcanic field by *Wilshire* and *McGuire* (1996), as well as in the oceanic crust of ODP Leg 176 (*Niu* et al., 2002). However, in the first case no massive Fe–Ti oxides are associated with melt pockets, although *Niu* et al. (2002) described them in the gabbros of South West Indian Ridge.

The range of variation of some incompatible element ratios of the gabbro xenoliths (e.g. Zr/Nb = 5-26; Y/Nb = 1.4-11), and particularly their *average* values (Zr/Nb = 9; Y/Nb = 3), indicate that these cumulates were derived from MORB-type liquids (*le Roux* et al., 2002). On the other hand these ratios are useless for rocks affected by significant hydrothermal metasomatism, like most of the studied diorites, since Cl-F-CO₂-bearing hydrothermal fluids are capable of mobilizing even those elements that are generally considered highly immobile (e.g. *Vard* and *William-Jones*, 1993; *Geisler* et al., 2002). In fact, the relatively high Nb contents in the diorites FB-h3 and FBY are positively correlated with CO₂ contents, and hence with the occurrence of secondary calcite in these rocks.

The origin of the diorites may be related to fractionation of residual liquids remaining after the formation of some of the cumulate gabbros since calcite-free diorites have Zr/Nb and Y/Nb signatures similar to those of gabbros. However the following additional hypotheses on the origin of diorites cannot be ruled out: (i) some of the Hyblean "diorites" may be derived from plagioclase-rich gabbro protoliths which have suffered a strong alkaline metasomatism; (ii) some diorite-related

liquids derive from small degrees of partial melting of gabbros during high temperature shearing (e.g. *Costa* and *Caby*, 2000).

Comparisons of trace element distributions between the xenoliths and the host rock exclude contamination as an important metasomatic agent for Hyblean xenoliths (Fig. 5). For instance, Sr and Ba content in most gabbros, related to plagioclase accumulation, exceeds that of the host rock. However an additional influx of these elements, i.e. interaction with percolating and uprising alkaline silicate melts (e.g. *Neumann* et al., 2000) cannot be excluded.

The positive correlation between LOI and LILE (see Table 6) and the presence of amphibole suggest that the concentration of these elements is also related to processes controlled by late-stage hydrous fluids. Hydrous fluids may also mobilise the alkalis and produce the Na–K-feldspar micro-veinlets cross-cutting the plagioclases of the diorites. Hydrous fluid circulation is in agreement with the relatively high $\delta^{18}O_{\text{SMOW}}$ values (FB12 = 8.7‰ and FB11 = 7.2‰). In fact, these values are similar to those of oceanic mafic rocks subjected to various processes including ocean floor weathering (*Staudigel* et al., 1981), but they also fall within the range of granulite xenoliths from deep continental crust (*Kempton* and *Harmon*, 1992).

According to *Scribano* (1988) and *Sapienza* and *Scribano* (2000), mafic granulites are of meta-igneous origin. The chemical compositions of these rocks suggest that they have suffered depletion during granulite-facies recrystallisation. The lack of major element correlation (Fig. 4) among gabbros, diorites and mafic granulites indicates that the latter may not derive from comagmatic protoliths.

Comparisons with gabbroids from different geodynamic settings

In order to define the nature of the Hyblean lower crust and the processes through which it may have evolved, the geochemical and petrographic features of the Hyblean lower crustal xenoliths were compared to igneous (cumulates in most cases) and metamorphic-textured gabbroids from well-known geodynamic settings.

The suite of Hyblean lower crust xenoliths consists of mafic granulites, subordinate gabbros and rare diorites. P–T data from the granoblastic mafic granulites indicates an intermediate pressure of metamorphism, suggesting the existence of a relatively thinned crust. The predominant mineral assemblage (two pyroxenes + plagioclase + Al-spinel) of the Hyblean mafic granulite xenoliths is analogous to that of the European Phanerozoic lower crust, though the latter often bear garnet (Downes, 1993). It is possible that the igneous protolith of the mafic granulites was formed by basaltic underplating of continental crust. However, petrographic evidence and geochemistry of the gabbros and diorites argue against a continental origin. Gabbros show shearing microstructures that are related to the melt pockets and Fe-Ti oxides III. Severe shearing and massive oxide patches within a silicate matrix are uncommon in gabbros from continental layered intrusions (Hunter, 1996). The continental occurrences of heavily sheared gabbros are rather unique (the Ivrea Verbano Zone, Northern Italy; *Rivalenti* et al., 1984 and Zingg, 1990). However, such modal and textural characteristics have been described in oceanic gabbroids, e.g. from transform segments of low-spreading ocean ridges (Vissers and Nicolas, 1995). These petrographic features are also very common in some ophiolitic ferrogabbros and ferrodiorites, i.e. those from the Italian Northern Apennines



Fig. 7. C1-normalised REE patterns and PM-normalised spiderdiagrams for gabbroids from different geodynamic settings. See text for references. Representative Hyblean xenoliths are reported: black line = average gabbro; dashed line/asterisk = average mafic granulite; field with dotted line = diorites

and Western Alps (*Pognante* et al., 1983; *Tribuzio* et al., 1995, 1999; *Tiepolo* et al., 1997).

Another argument supporting the oceanic affinity of Hyblean gabbros is the covariation between the anorthite content of plagioclase and the Mg# value of coexisting clinopyroxene (Fig. 3). Most data for the Hyblean oxide-rich gabbros is in agreement with the fields defined by oceanic gabbro xenoliths from the Canary Islands (*Schmincke* et al., 1998), while the oxide-poor gabbros fall within the range of Mid Ocean Ridge cumulates (*Ross* and *Elthon*, 1993: Fig. 3).

In Fig. 4, the whole-rock major element composition of the Hyblean gabbros is compared with that of oxide-rich gabbros from the South West Indian Ridge (oceanic crust; *Coogan* et al., 2001), gabbroids from Platta (ocean-continent transition; *Desmurs* et al., 2002) and Calabria (continental; *Morten*, unpublished data). The suite of Hyblean lower crustal xenolithst mostly resembles those of the Platta gabbroic suite (including Mg-, Fe-, Fe–Ti–P-gabbros and diorites) and the oxide-gabbros from South West Indian Ridge.

Figure 7 shows C1-normalized REE patterns and N-MORB-normalized spiderdiagrams for gabbroids (cumulates in some cases) from different geodynamic settings. The enrichment of LREE over HREE is considered typical of the lower continental crust ($La_N/Yb_N > 1$; *Downes*, 1993). Nevertheless some exceptions exist (e.g. metagabbros from Calabria, Pannonian Basin). Conversely, gabbroids with oceanic affinity usually show almost flat or LREE depleted patterns ($La_N/Yb_N < 1$; *Coogan* et al., 2001). The Hyblean diorites show REE patterns and N-MORB-normalised spiderdiagrams similar to the plagioclase-rich mafic granulites from Queensland, although the latter have lower LREE and trace element concentrations. *Rudnick* (1992) interpreted such REE patterns as conforming to those of plagioclase in equilibrium with mafic to intermediate liquids. The N-MORBnormalised trace element distribution of Hyblean gabbros is also comparable to those of the West Iberian Margin (*Cornen* et al., 1999), Pannonian Basin (*Embey-Isztin* et al., 2003) and oxide-rich gabbros from South West Indian Ridge (*Coogan* et al., 2001), with the main differences regarding the Sr and Ba anomalies.



Fig. 8. Relations between Rb/Nb and K/Rb ratios. Same symbols as in Fig. 4. Open star = Lower Continental Crust (*Weaver* and *Tarney*, 1984); black star = LOR (host basalt); grey star = N-MORB (*Hoffmann*, 1988); full circle = P-MORB (*le Roex*, 1987). For comparison, the fields of gabbroids from different geodynamic settings are also shown (see text for references)

The Hyblean lower crustal xenoliths and the gabbroic suites used as comparison are plotted in Fig. 8 (K/Rb *vs* Rb/Nb binary plot). Continental gabbroids exhibit higher K/Rb and lower Rb/Nb ratios than the oceanic and ocean-continent transition gabbroids. The Hyblean xenoliths partially overlap the fields of granulites from the Pannonian Basin and gabbroids from Canary Islands and South West Indian Ridge. Conversely, they lie far from the representative points of continental crustal gabbroids and lower continental crust.

Concluding remarks

The Hyblean xenolith suite consists of upper mantle lithologies, lower-crustal mafic granulites, minor oxide-poor and oxide-rich gabbros, rare diorites, and samples from different levels of the sedimentary sequence at time of the eruption (Upper Miocene). Felsic and garnet-bearing lithotypes are absent from the suite. The pet-rological and geochemical data discussed above suggest the following conclusions:

- 1. The Hyblean gabbros are cumulates formed from liquids with MORB affinity. Strongly fractionated Fe–Ti-rich melts of probable intercumulus origin have been filter-pressed and then injected into adjacent gabbroic masses during shearing. These injected liquids precipitated oxide minerals, giving rise to the oxide-rich gabbros.
- 2. Some gabbros and most diorites show important chemical and mineralogical transformations (i.e. pervasive alkali metasomatism of plagioclase), related to late-stage and hydrothermal fluids which caused LREE and alkali enrichments.
- 3. Mafic granulite xenoliths are not comagmatic with the other xenolith groups. Hence they are almost unaffected by metasomatism and hydrothermal alteration and probably derived from the lowest section of the crustal column.
- 4. The petrographic features of studied oxide-rich gabbros closely resemble those of gabbros drilled and recovered in oceanic environments, especially in the fracture zones of slow-spreading ridges.

These petrographic considerations on the crustal xenoliths can be combined with the following evidence: (i) geophysical investigations (*Arisi Rota* and *Fichera*, 1987; *Della Vedova* et al., 1989) indicating the presence of a thick mafic (likely gabbroic) body just underneath the sedimentary succession; (ii) logs from commercial boreholes having recovered gabbro and peridotite fragments in the deepest parts of the Gela Nappe (*Longaretti*, 1986, personal communication: see Fig. 9); (iii) oil wells in the Ragusa area, south-western part of the Hyblean Plateau, having reached tholeiite gabbros and dolerites underneath Triassic dolomite, with no evidence of pyrometamorphism, about 2500 m below sea level (*Cristofolini*, 1966 and therein references).

The above documentation suggests that the Hyblean sedimentary succession stands upon gabbroic bodies. Thus, it appears improbable that the ascending magma has sampled the bottom (upper mantle, granulitic lower crust) and top (gabbro, sedimentary sequence) of the lithospheric column, but left the middle untouched. Hence we can reasonably assume that the studied xenoliths adequately represent the Hyblean crustal basement, which may consist of a "fossil" peridotite-gabbro ridge (Fig. 9). Nevertheless, in case the above assumption is not agreed, the continental nature of the Hyblean lithosphere cannot be definitively ruled out.



Fig. 9. Hypothetical NW–SE profile of the crustal structure across the Hyblean Plateau and neighbouring areas, according to the conclusions of the present study. Heights are exaggerated for clarity. All parts of this xenolith-based profile are merely conceptual and not to scale

Although the petrology of the xenoliths fits either an oceanic or oceancontinent-transition (OCT) nature for the peridotite/gabbro ridge, the oceanic hypothesis is more appropriate, because the xenolith-bearing diatremes are located well within the Plateau, up to 35 km from the Malta escarpment, on its land-side (Fig. 1). A supposed OCT section would, instead, have been placed on the sea-side of the slope, like the present-day Galicia and Gorringe Banks, West Iberia Margin (*Cornen* et al., 1999).

The conclusion that the lower crust beneath the Hyblean Plateau may not belong to the African continental crust, as commonly considered in the literature (*Burollet* et al., 1978), could have important implications for the geological setting of this area and, more generally, this hypothesis raises further questions concerning the geodynamic reconstruction of the Central Mediterranean area. The Hyblean lower crust may represent, like the adjacent Ionian crust, a relic of the Permo-Triassic Palaeo-Tethys ocean (*Vai*, 1994). On these grounds, the Malta escarpment, generally considered an ancient passive continental margin (*Finetti*, 1982), may instead represent a "fossil" segment of a transform margin.

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V. Scribano et al.: Gabbroic xenoliths from the Hyblean Plateau

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