THE ANTRONA NAPPE: LITHOSTRATIGRAPHY AND METAMORPHIC EVOLUTION OF OPHIOLITES IN THE ANTRONA VALLEY (PENNINE ALPS)

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ABSTRACT

The Antrona ophiolite (western Central Alps) represents a tectonic fragment of the oceanic lithosphere of the Upper Jurassic - Lower Cretaceous Ligurian-Piedmont basin, a section of the Western Alpine Tethyan Ocean. The Antrona ophiolite occurs at lower structural levels in the Alpine nappe stack and is sandwiched between the overlying continental Monte Rosa Nappe (upper Penninic) and the underlying Camughera-Moncucco continental Unit (middle Penninic). The Monte Rosa Nappe is overlain by the Zermatt Saas ophiolitic Unit. Despite the tectono-metamorphic reworking, the Antrona ophiolite exhibits all typical lithologies of oceanic lithosphere: ultramafic and mafic plutonic rocks, mafic volcanic rocks and deep-sea sediments can be recognized. Several rock types were distinguished among mafic rocks. New findings and inferences are: i) the occurrence of probable relics of magmatic structures in some amphibolites, inferred to be pillow lavas or pillow breccia; ii) the occurrence of lawsonite pseudomorphs-bearing amphibolites, not described so far in the study area as precursor of the origin of epidote-amphibolites the Antrona Valley. A qualitative P-T diagram deduced from the stability conditions of mineral parageneses is presented and compared with published P-T paths for the Antrona and Zermatt-Saas ophiolites. The metamorphic evolution of the studied rocks is characterized by blueschite prograde path followed by high pressure (eclogitic) metamorphic peak. P-T estimates for the metamorphic peak were calculated by the Na-clinopyroxene garnet equilibria and the jadeite content in omphacite. T = 372° C for a nominal pressure of P = 1 GPa and T = 386° C for a nominal pressure of P = 1.5 Gpa were obtained. Jd₃₀ as maximum Jadeite content suggests P > 1 Gpa. Retrograde path, although not well constrained, is dominated by epidote-amphibolite/amphibolite facies conditions, in accord with published data, differing from those inferred so far for the overlying Zermatt-Saas ophiolite.

INTRODUCTION

The Zermatt-Saas and Antrona metaophiolites represent remnants of the oceanic lithosphere of the Mesozoic western Tethys, now dismembered as tectonic slices in the internal Pennine domain of the western Central Alps. The Antrona Unit obtained not much attention in the past, despite its crucial tectonic position within the nappe stack.

This paper presents the results of a geological study carried out on the eastern side of the Antrona Valley, a tributary of the Ossola Valley (western Central Alps). Despite the polyphase metamorphic history, the Antrona ophiolite exhibits all typical lithologies of oceanic lithosphere, i.e., ultramafic and mafic plutonic rocks, mafic volcanic rocks and deep-sea sediments. This work focuses on plutonic and volcanic metabasites. Among the mafic volcanic rocks (commonly described in literature as "amphibolites") several lithologies, often retaining relict magmatic structures, were distinguished. Lozenge-shaped porhyroblasts consisting of epidote+chlorite aggregates were observed for the first time in amphibolites from the study area. They are interpreted as pseudomorphs of former lawsonite, thus providing a clue to discuss the metamorphic evolution of the Antrona ophiolite. Temperature and pressure conditions were estimated for the high pressure metamorphic peak, whereas the prograde and retrograde P-T diagram was deduced from the stability conditions of mineral parageneses.

GEOLOGICAL SETTING

Regional geology

In the Pennine-Lepontine nappe stack (e.g., Escher et al., 1987; 1993), the Zermatt-Saas ophiolite nappe mantles the

Monte Rosa Unit and disappears below the Mischabel backfold; the Antrona ophiolite is situated on the footwall of the Monte Rosa Nappe (Fig. 1) and in turn overlays the middle Penninic Camughera-Moncucco Nappe (Bearth, 1956; Laduron, 1976; Bigioggero et al., 1981; Keller et al., 2005b). The Antrona and the Zermatt-Saas metaophiolites almost completely envelop the Monte Rosa Nappe, with the exception of the area between Gornergrat and "Passo della Preja", where the Furgg Zone is dominant and only thin slices of ophiolite rocks are present (Fig. 1).

The Monte Rosa Nappe is a large NW-vergent recumbent anticline, strongly refolded by S-vergent backfolds of supposed Oligocene age (Milnes et al., 1981; Escher et al., 1997; Steck et al., 1997). The Monte Rosa Nappe, together with the Gran Paradiso and Dora Maira "massifs", constitute the inner and upper continental nappe system of the Penninic Zone in the Western Alps. In the Monte Rosa Nappe, two main pre-Alpine protoliths have been recognized: a pre-granitic metamorphic complex (Bearth, 1952; Gosso et al., 1979; Dal Piaz, 1993; Keller and Schmid, 2001; Keller et al., 2005a), consisting of high grade paragneiss rich in mid- to coarse-grained pegmatites, and a granitic complex represented by dominant granitic-granodioritic bodies cut by aplitic and pegmatitic dykes of Late Carboniferous and Permian ages (Hunziker, 1970; Frey et al., 1976). During the Alpine orogeny, the pre-granitic complex has been reworked into garnet micaschists and albitic schists, and the granitic bodies into massive, schistose or mylonitic orthogneiss.

In the study area, the Antrona ophiolite is separated from the Monte Rosa basement by the Furgg Zone (Argand's "synclinal de Furggen", Argand, 1911). The Furgg Zone consists of micaschists, albitic schists, and leucocratic gneisses (Permian-Carboniferous?) with eclogitic to greenschist-facies mafic boudins, thin micaceous quartzites and dolomitic mar-

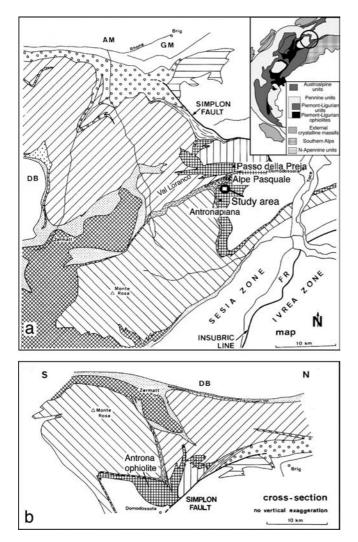


Fig. 1 - Tectonic map (a) and cross-section (b) of the Pennine Alps in the western Central Alps (modified after Milnes et al., 1981). The inset shows the location of the study area in the central-western portion of the Alpine chain.

ble of pre-Mesozoic (?) age (Bearth, 1954). In the upper part, it includes layers (tectonic transposition) of Mesozoic metasediments (tabular quartzites, dolostone, carnieule, calcschists) and tectonically transposed ophiolite rocks of the underlying Antrona Unit (Blumenthal, 1953; Bearth, 1957; 1964; Dal Piaz, 1966; Wetzel, 1972;). According to Dal Piaz (Dal Piaz, 1964; 1966), the Furgg Zone exposed in the southern Monte Rosa has a pre-granitic protolith, free of Mesozoic rocks and was affected by eclogitic to greenschists facies Alpine metamorphism. More recently, the Furgg Zone has been interpreted as cover portions of the Monte Rosa basement (Jaboyedoff et al., 1996; Escher et al., 1997; Steck et al., 2001) or of the continental Portjengrat Unit (Keller and Schmid, 2001). Alternatively, the Furgg Zone has been considered as the suture of the Valais basin (Froitzheim, 2001), or a tectonic mélange interposed between the continental Monte Rosa-Portjengrat Units and the Zermatt-Saas and Antrona ophiolites (Kramer, 2000). Based on SHRIMP analyses, Liati et al. (2001) interpreted the Furgg Zone as a tectonic mélange with a strongly heterogeneous composition including Variscan and pre-Variscan basement (510 \pm 5 Ma) as well as post Variscan rocks (272 ± 4 Ma), which were affected by Alpine metamorphism (87 \pm 20 Ma is the age of metamorphism determined in an amphibolitized eclogite boudin from the Andolla area in the upper Antrona Valley).

The Zermatt-Saas and Antrona metaophiolites, the Monte Rosa Nappe, the Furgg Zone, and the Camughera Moncucco Unit have all experienced regional subduction-related highpressure metamorphism during the Alpine orogeny. The Zermatt-Saas ophiolite subduction history is attested by the occurrence of lozenge-shaped pesudomorphs of lawsonite and other prograde relics followed by peak eclogitic to UHP mineral assemblages (Ernst and Dal Piaz, 1978; Oberhänsli, 1980; Barnicoat and Fry, 1986; Martin and Tartarotti, 1989; Reinecke, 1991; 1998). Eclogite rocks in the Antrona ophiolite have been described by Colombi and Pfeifer (1986). In the Monte Rosa Nappe, the Alpine eclogitic imprint has been recognized in both the pre-granitic complex (Dal Piaz and Lombardo, 1986; Borghi et al., 1996; Michard et al., 1996; Keller et al., 2005a) and in the orthogneisses (Dal Piaz and Gatto, 1963; Dal Piaz and Lombardo, 1986). More recently, eclogite mineral assemblages have been recognized also in the Camughera-Moncucco Nappe (Keller et al., 2005b).

During decompression accompanying exhumation, the Zermatt-Saas and upper Monte Rosa Units were re-equilibrated under greenschist facies conditions (Frey et al., 1974; Ernst and Dal Piaz, 1978; Keller et al., 2005a). The lower parts of Monte Rosa and the Antrona ophiolite underwent epidote-amphibolite facies metamorphism (Pfeiffer et al., 1989).

The Antrona Nappe

The Antrona Nappe is exposed in the Antrona-Loranco-Anzasca Valleys, on the Italian side, and in the Laggintal -Simplon area, on the Swiss side (Fig. 1). In Italy, the Antrona ophiolites are also exposed in the Bognanco Valley (north-east of Antrona) and can be followed to the Vigezzo Valley east of Domodossola. In the lower Bognanco Valley, rocks interpreted as belonging to the Antrona ophiolite define the contact between the Camughera-Moncucco and Monte Leone Nappe (Keller et al., 2005b). The main lithologies in the Antrona area (Antrona and Anzasca Valleys) were described by Laduron and Merlin (1974), Laduron (1976), Colombi and Pfeifer (1986), Colombi (1989), Pfeifer et al. (1989). The typical Antrona ophiolite suite includes serpentinized ultramafites, metagabbros and metabasic rocks. In the nearby Bognanco Valley, Carrupt and Schlup (1998) described a detailed ophiolitic section consisting of N-MORB to T-MORB-type metabasalts and minor serpentinite and gabbros. This ophiolite is covered by (from bottom to top) garnet-micaschists, metaquartzites, calcschists and intercalated metabasite, sedimentary breccia, pure calcschists, marbles, and graphitic marbles. The Antrona ophiolite is dominated by a Tertiary amphibolite facies metamorphism. Mafic eclogites containing garnet, symplectites, amphibole and relict omphacitic clinopyroxene have been described by Colombi and Pfeifer (1986) in the Antrona-Anzasca area. Minimum pressures of 14-16 kbars at 480-720 °C were suggested for the high pressure stage of these rocks by these authors.

FIELD RELATIONSHIPS OF OPHIOLITE ROCKS IN THE ANTRONA VALLEY

The study area is located in the upper Antrona Valley, between Alpe Pasquale, to the north, and the village of Antronapiana, to the south (Fig. 1); it was mapped in detail

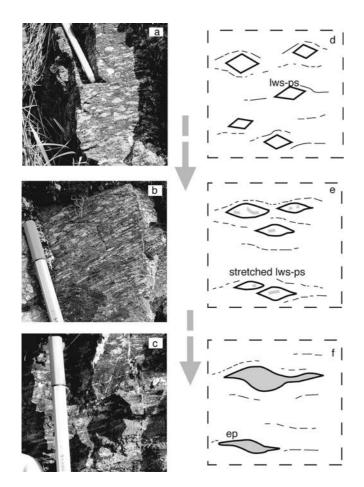


Fig. 2 - Photographs of lawsonite (pseudomorphed)-bearing amphibolites (a, b) and epidote-amphibolite (c). d-e-f) Cartoons showing the microstructural evolution and mineral changes from lawsonite-bearing amphibolite to epidote-amphibolite. a, d) Lozenge-shaped pseudomorphs (lws-ps) consisting of epidote, chlorite, and carbonate. b, e) Pseudomorphs are deformed by stretching. c, f) Stretched pseudomorphs evolve to mm-thick layers, mostly consisting of epidote.

at the scale of 1: 5000 by Turco (2004). In this area, metamorphosed ophiolites are exposed along the eastern side of the Antrona Valley. The main rock types are ultrabasic rocks, metagabbros and metabasalts. Ultrabasic rocks are mostly serpentinized and make up a 35 km² body extended from Antronapiana to Pizzo Ciapé, near the tectonic contact with the Camughera-Moncucco Unit. Serpentinite structures range from massive to mylonitic. Mylonitic serpentinites are interlayered within metabasites. Dark green fine-grained amphibolite dikes are interlayered within massive serpentinites near Antronapiana. The contact between serpentinites and metabasalts at the northern border of the massif is commonly marked by actinolite-schists. To the south of Antronapiana, serpentinites are in tectonic contact with Mgmetagabbros. This contact does not show any reaction zone. Outcrops of metagabbros are rare and restricted to m-scale layers associated with metabasalts. A body of Mg-metagabbro up to 50 m in thickness crops out at Pizzo Ciapé, near the divide with the Bognanco Valley. Metagabbro has transitional contacts with prasinite and their macroscopic distinction is usually based on structure and grain size .

Metabasites are the most abundant rock type in the studied ophiolite suite. These rocks are characterized by fine grain-size and dark blackish-green colours. On the basis of mineral components and structure, we differentiated the fol-

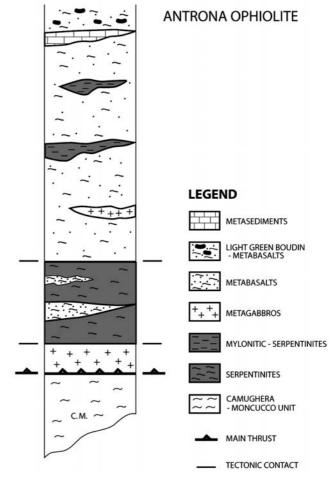


Fig. 3 - Simplified lithostratigraphic column of the ophiolite suite in the Antrona valley.

lowing types: amphibole-eclogites, garnet-amphibolites, lawsonite (pseudomorphosed)-bearing amphibolites, epidote- amphibolite, pale green microboudin-bearing amphibolites. Amphibole-eclogites are very scarce (one main mscale layer near the Fornalino Pass). The high-pressure mineral assemblage consists of garnet-Na-clinopyroxene-rutile and amphibole \pm quartz.

Amphibolites are the dominant rock type in the working area. Garnet-rich amphibolites are fine-grained dark green rocks characterized by the strong mineral lineation of amphibole crystals. Lawsonite (pseudomorphosed)-bearing amphibolites (see next section) are macroscopically recognizable for the occurrence of whitish, centimetric lozengeshaped crystals (Fig. 2). So far, lawsonite (now pseudomorphosed)-bearing rocks have not been documented in the study area. Epidote-amphibolites are typically characterized by the presence of millimetric yellow-green epidote-rich layers alternating with dark green amphibole-rich layers. One rock type commonly grades into the other (Figs. 2a, 2c). The structural evolution from one type to the other is marked by increasing stretching of the cm-sized and lozenge-shaped pseudomorphs on lawsonite, followed by crystallization of epidote in planar, mm-thick layers (Figs. 2d, 2f). A further rock type recognized in the field (best outcrops along the right side of the Alpe Cavalli Lake in the

Table 1 - Lithologies and	netrographic features	of the studied metao	phiolite in the Antrona valley.
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Rock	Micro-structure	Eo-Alpine mineralogy	Meso- and Neo-Alpine mineralogy
Metagabbro	Flaser to mylonitic	(Na-clinopyroxene? zoisite?), amphibole	Amphibole, chlorite, clinozoisite, pistacite
Amphibole Eclogite	Massive polygonal to interlobate	Garnet, Na-clinopyroxene, amphibole core, rutile	Amphibole rim, amph-plag symplectites, titanite
Garnet amphibolite	Nematoblastic-porphyroblastic	Garnet, amphibole, feldspar, quartz	Chlorite, clinozoisite, amphibole
Lawsonite -bearing amphibolite	Nematoblastic	(Laws), amphibole, garnet, rutile	Amphibole, chlorite, biotite, plag, epidote, titanite
Epidote amphibolite	Nematoblastic	Amphibole, (laws?, garnet?),	Pistacite, chlorite, amphibole
Microboudin-bearing amphibolite	Nematoblastic- symplectitic?	Amphibole, garnet, (Na- clinopyroxene?), rutile	Amphibole-plag symplectite, amphibole around garnet, clinozoisite
Prasinite	Granoblastic to schistose	no relics	Amphibole (Actinolite), plag, chlorite, epidote, quartz
Serpentinite	Porphyroclastic to schistose to mylonitic	Olivine, serpentine, spinel	Serpentine, magnetite
Quartzite	Granoblastic to schistose	Quartz, white mica, garnet	Quartz, white mica, brown biotite
Micaschist	Schistose-porphyroblastic	White mica, garnet, chloritoid, quartz	White mica, brown biotite, chlorite
Marble	Granoblastic	Carbonate, quartz, biotite, white mica	Chlorite, epidote
Calcschist	Schistose-granoblastic	Carbonate, quartz, white mica,	Clinozoisite, biotite

Loranco Valley) is characterized by the occurrence of dmscale boudins of pale green amphibolite inside dark green amphibolite. Lawsonite pseudomorphs, epidote-rich layers and boudins within amphibolites show a preferential orientation parallel to the regional main schistosity.

Metasediments of the ophiolite suite in the Antrona Valley are mostly represented by garnet-quartzites, micaschists, marbles and calcschists.

A representative lithostratigraphic column of the ophiolite suite mapped in the Antrona Valley is shown in figure 3.

PETROGRAPHY

Rock types and microstructural features of the investigated samples are summarized in Table 1. In the studied rocks four main foliations were recognized, related to ductile deformation phases D1-D4. S1 is the oldest foliation, locally preserved inside garnet; S2 corresponds to the regional foliation seen in the field, and is mostly marked by amphibole and garnet; S3 and S4 are younger foliations cutting S2, and are commonly defined as crenulation cleavage or S-C structures marked by chlorite and albite. All ductile structures are cut by veins filled with prehnite-pumpellyite facies minerals. Four stages of mineral crystallization (B1-B4) were recognized in the studied mafic rocks (see Fig. 9 and Discussion).

Ultrabasic rocks

Ultrabasic rocks consist largely of serpentine (antigorite), magnetite, and minor diopside and talc. Olivine can be found in more massive rock types, where it is present in coarser-grained (mm) stretched grains (olvine I) and finegrained neoblast (olivine II). This texture suggests that olivine I underwent grain-size reduction by recrystallization during the Alpine deformation (Fig. 4a). Coarsegrained olivine exhibits undulatory extinction and subgrains. The shape of subgrains reminds the microstructure of porphyroclastic olivine in mantle peridotites (Mercier and Nicolas, 1975; Nicolas and Poirier, 1976). Amphibolite dikes interlayered with massive olivine-bearing serpentinites consist of amphibole (85-90%), plagioclase (5-10%), titanite (5%), epidote (1%). They exhibit a fine-grained granoblastic texture.

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Metagabbro

Metagabbro samples consist of amphibole (50-60%), epidote (15-30%), plagioclase (10-20%), quartz (5-10%), chlorite (2-10%), opaque (1%), \pm biotite, carbonate. Their microstructure varies from weakly schistose with relics of magmatic microstructures, to flaser and mylonitic. In flaser metagabbro, the foliation (S2) is spaced and defined by amphibole films alternating with lithons made of epidote and plagioclase (Fig. 4b). Amphibole, epidote and plagioclase crystallization pertains to B3 stage. Foliation wraps around mm-sized, sigmoidal or stretched porphyroclasts of former clinopyroxene now replaced by coarse-grained amphibole grains. Strain shadows are commonly filled with fine-grained amphibole, quartz, \pm epidote, chlorite (stages B3, B4).

Amphibole-eclogite

This rock consists of medium-grained, inequigranular garnet (35-40%), amphibole (20-45%), Na-clinopyroxene (10-20%), amphibole + plagioclase symplectites (2-10%), \pm quartz, rutile (Fig. 4c). Microstructure ranges from polygonal to interlobate. Garnet is present in subhedral to euhedral

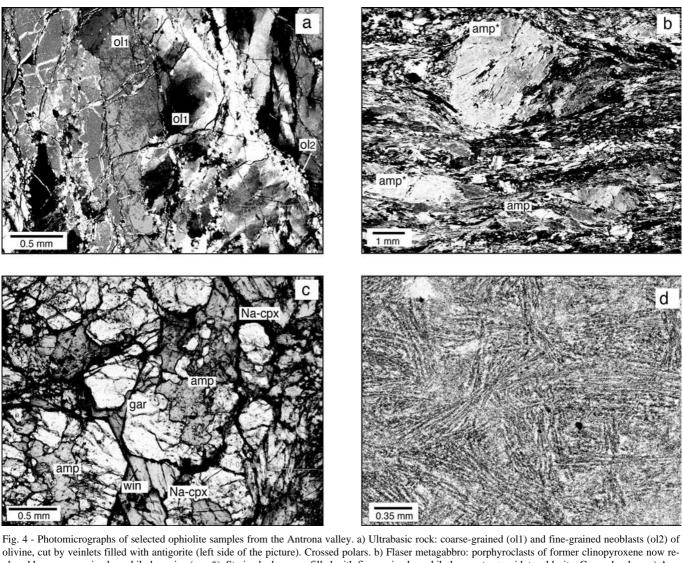


Fig. 4 - Photomicrographs of selected ophiolite samples from the Antrona valley. a) Ultrabasic rock: coarse-grained (ol1) and fine-grained neoblasts (ol2) of olivine, cut by veinlets filled with antigorite (left side of the picture). Crossed polars. b) Flaser metagabbro: porphyroclasts of former clinopyroxene now replaced by coarse-grained amphibole grains (amp*). Strain shadows are filled with fine-grained amphibole, quartz, ± epidote, chlorite. Crossed polars. c) Amphibole-bearing eclogite: garnet (gar), partially replaced by subhedral Ca-amphibole (amp), is in contact with both winchite-rich amphibole (win) and Na-clinopyroxene (Na-cpx). Plane polarised light. d) Microboudin-bearing amphibolites: aggregates of acicular crystals with "dendritic"-like or "sheaf"-like structure occurring in a microboudin (see also backscattered picture in Fig. 6b). Plane polarised light.

porphyroblasts or poikiloblasts (scattered inclusions of amphibole). Amphibole occurs either as nematoblasts in textural equilibrium with both garnet and Na-clinopyroxene, or subhedral grains replacing garnet. Amphibole nematoblasts commonly have uncoloured cores and blue-green rims, which correspond to chemical zoning (see next section and Fig. 6b). The core of these amphibole crystals, together with garnet and Na-clinopyroxene, are inferred to pertain to the HP mineral paragenesis (stage B2). Na-clinopyroxene occurs as greenish nematoblasts showing textural equilibrium with garnet and amphibole. The contact between garnet and clinopyroxene is almost always marked by a very thin (um scale) rim of green amphibole (stage B3) as a reaction product of the two HP mineral phases. Amphibole + plagioclase symplectites were produced by the break-down of Naclinopyroxene, reacting with quartz to form amphibole + sodic plagioclase through hydration reactions with adjacent minerals during retrogression.

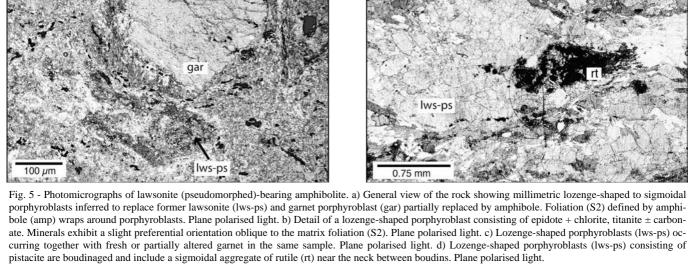
Garnet-amphibolite

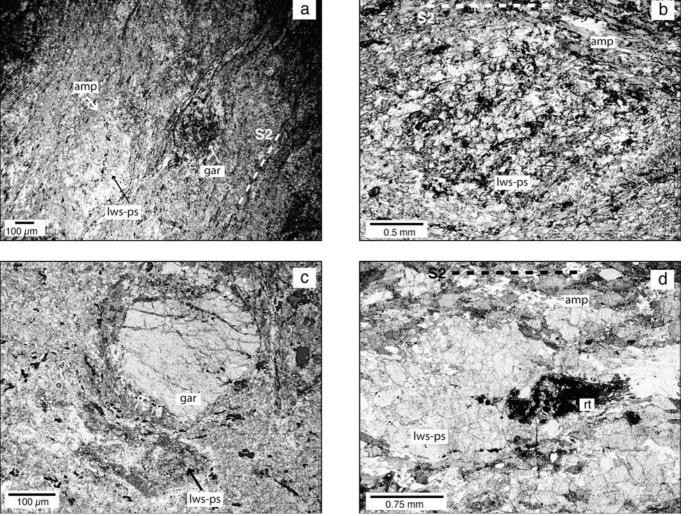
The rock consists of amphibole (30-45%), plagioclase

(10-25%), epidote (18-20%), carbonate (0-10%), white mica (0-10%), opaque minerals (3-5%), garnet (1-2%), \pm quartz, chlorite, titanite. Garnet poikiloblasts are up to 5mm in diameter and include epidote, amphibole, opaque minerals. Garnet may be either partially replaced by green amphibole at the rim of porphyroblasts, or completely overgrown by breakdown products such as amphibole, epidote, plagioclase, and opaque minerals, still retaining the shape of the original mineral. The foliation (S2) of the rock is defined by shape preferred orientation (SPO) of green or blue-green amphibole and epidote (stage B3).

Lawsonite (pseudomorphed)-bearing amphibolite

The rock consists of amphibole (40-60%), epidote (20-30%), plagioclase (15-25%), garnet, chlorite (5%), carbonate (0-5%), rutile-titanite (1%), \pm opaque, biotite, quartz. The main foliation (S2) is defined by amphibole SPO (stage B3). Millimetric lozenge-shaped porphyroblasts made up of epidote (clinozoisite and pistacite) + chlorite, titanite \pm carbonate aggregates were identified (stages B3-B4). The shape of these porphyroblasts range from lozenge to sig-





moidal, according to the intensity of shearing affecting the rock (Figs. 5a, 5b). In general, the porphyroblasts show a preferential shape orientation; their forming minerals exhibit a slight preferential orientation almost oblique to the matrix foliation (Fig. 5b). Lozenge-shaped porphyroblasts may occur together with fresh or partially altered garnet in the same sample (stage B2?; Fig. 5c). In some samples the lozenge-shaped aggregates are deformed by boudinage or are stretched to form mm-thick layers parallel to the foliation. In one sample, microboudins consisting of pistacite include a sigmoidal aggregate of rutile near the neck of boudins (Fig. 5d). This texture suggests that lawsonite and rutile coexisted before or during boudinage affecting the foliation (S2) defined by epidote + amphibole.

In all samples, the matrix foliation wraps around the lozenge-shaped porphyroblasts, as well as garnet porphyroblasts (see Figs. 5a) indicating that both porphyroblasts growth took place before or during the ductile deformation generating the foliation. Based on the shape of porphyroblasts and on their forming minerals, we interpret the porphyroblasts as pseudomorphs after former lawsonite. Epidote + chlorite represent the main products of lawsonite breakdown.

Epidote amphibolite

The common mineral assemblage includes epidote (40-60%), amphibole (20%) plagioclase (5-15%), carbonate (0-10%), chlorite (5-20%), brown biotite (5-10%), opaque minerals (1-12%), titanite (1-2%), and quartz (1-2%). Foliation (S2) is defined by compositional layering marked by mm- to cm epidote-rich layers alternating with amphibole-rich layers, and by SPO of epidote and amphibole crystals (stage B3). In one sample, S2 foliation marked by green amphibole is cut by mm-scale veins filled with prehnite (later than B4).

Microboudin-bearing amphibolites

The host rock consists of amphibole (35-50%), microcrystalline plagioclase + amphibole symplectites (20-30%), plagioclase (15-29%), epidote s.l. (20-25%, carbonate (0-5%, opaque minerals (ca. 1%) \pm garnet, white mica, rutile. Microstructure is characterized by mm- to cm compositional layering of symplectites (undetermined under the microscope) alternating with layers of amphibole + epidote + plagioclase. Boudins (cm-scale) have a microcrystalline symplectitic structure, similar to that of symplectite layers in the host rock. Some portions of symplectites in the boudin consist of aggregates of acicular crystals arranged in a "dendritic"-like or "sheaf"-like structure (see Fig. 4d; see also backscattered picture in Fig. 6a). It is inferred that microcrystalline symplectites (stage B3) correspond to the breakdown of former Na-clinopyroxene, whilst aggregates with "sheaf"-like structure are due to recrystallization of symplectites (B4?). Petrographic and microstructural analysis suggest that the microboudin-bearing amphibolites represent transposed micropillow or pillow-breccia basalts.

MINERAL CHEMISTRY AND THERMOBAROMETRY

Analytical methods

Mineral analyses of selected samples of the Antrona

ophiolite rocks were performed at CNR-IPDA in Milano by using a SEM Cambridge System Stereoscan 360, equipped with an Energy Dispersive Spectrometer (EDS). The correction program is ZAF4 (Pouchou and Pichoir, 1985). Operating conditions were: accelerating voltage = 20kV, working distance = 25 mm, probe beam = 400 picoampere, count time = 50s. Selected SEM-EDS analyses are reported in Table 2.

Only eclogite samples were analysed also at CNR-IGG in Padova with a Cameca SX50 electron microprobe, equipped with four WDS spectrometers. Operating conditions: 15 KeV acceleration voltage and 15 nA sample current; count time = 10 sec at peak, 5 sec at background. The correction program is PAP (Kato, 2005). Natural and synthetic standards were used.

SEM-EDS analyses

SEM-EDS technique is particularly useful for revealing detailed microstructures, such as symplectites or intergrowths, chemical heterogeneity, etc., providing at the same time (although semi-quantitative) chemical compositions of the observed mineral phases (e.g., Vernon, 2004 and refs. therein). In this study, SEM-EDS was utilized for analyzing the chemical heterogeneity in eclogite amphibole and the fine-grained "symplectites" in the pale green microboudins occurring in amphibolites (see photomicrographs in Figs. 4c and 4d, respectively).

Backscattered images revealed chemical heterogeneities of amphiboles in eclogite (Fig. 6a). SEM chemical compositions of amphibole in eclogites are plotted in Fig. 7; selected analyses are reported in Table 2. Amphibole in the eclogite groundmass is zoned with a core of winchite and rim of Mgkatophorite, taramite, and Fe-tschermakite (Figs. 7d, 7c and 7b, respectively). Amphibole coronas in contact with garnet range from barroisite to Fe-pargasite and Fe-tschermakite (Figs. 7d, 7a). In eclogites, amphibole is also included in garnet; its composition ranges from actinolite and Fe-pargasite to barroisite (Figs. 7a, 7b, 7d).

Backscattered images show that microboudins in amphibolites are characterized by granoblastic structure (Fig. 6b). Mineral phases constituting this fine-grained rock are actinolite, clinozoisite-pistacite, and albite (Table 2).

Representative compositions of amphibole crystals from amphibolites and metagabbro are plotted in Fig. 8. A selection of these analyses is reported in Table 2. Most amphiboles are Ca-amphiboles, ranging in composition from actinolite to Mg-hornblende (Figs. 8a, 8d), and from edenite to Fe-pargasite (Fig. 8b). Barroisitic amphibole was found in almost all amphibolite samples (Fig. 8c).

Plagioclase of all the analyzed amphibolites and metagabbro shows a wide composition ranging from Ab = 70-99%, An = 1-30%.

Microprobe analyses of eclogite minerals

Mineral compositions of amphibole, clinopyroxene, and garnet from amphibole-eclogites are reported in Tables 3 and 4. These quantitative analyses were used for thermometry calculation (see next paragraph).

Amphibole in the groundmass of eclogite samples is not homogeneous. Chemical zoning from core to rim revealed by backscattered images (see Fig. 6a) and SEM-EDS analyses seems to be not confirmed by electron microprobe analyses that provide either actinolite or winchite compos-

						Α	AMPHIBOLE	LE						EF	EPIDOTE					PL	PLAGIOCLASE	SE	
Lithology	Eclogite	Eclogite	Eclogite Eclogite	Eclogite	Eclogite 1	Eclogite	Anf bd	Met gb	Met gb	Anf lw	Anf gr	Anf ep I	Lithology Amph bd Amph bd Amph ep Amph ep	nph bd Aı	nph bd Ai	nph ep Ai	nph ep	Lithology Amph bd Amph bd Amph lw Amph ep	nph bd An	nh bd An	nph lw Ar		Met gb
Sample	CI	CI	C14B	CI	CI	CI	C30	C49	C49	C52	C25	C26 5	Sample	C39	C39 C	C45AA C	C45AA	Sample	C30	C30	C54 C	C45AA	C55B
Analysis	113	114	144	107	111	112	27	5	6	68	22	138 /	Analysis	135	138	27	28	Analysis	49	60	229	36	92*
Occurrence	core	rim	cn	cu	incl	incl	sym			cu		-	Occurrence			core	rim						
SiO2 wt%	54.90	48.46	50.45	38.92	54.99	51.11	53.73	55.51	53.21	39.95	50.5	48.07	SiO2 wt9	38.79	38.26	37.60	38.37	SiO2 wt%	60.48	67.86	63.56	62.61	60.05
TiO2	0.14	0.09	0.16	0.02	0.05	0.16	0.01	0.01	0.1	0.57	0,00	0.29	Ti02	0.00	0.18	0.11	0.13	Ti02	0.00	0.00	0.06	0.09	0.08
A12O3	3.27	9.25	8.32	17.52	1.79	7.94	2.1	2.2	5.59	17.66	8.31	7.59	A12O3	29.45	28.32	23.48	28.56	A12O3	24.8	19.11	23.87	22.41	24.55
MnO	0.15	0.15	0.22	0.2	0,00	0.1	0.03	0.1	0.1	0.16	0.3	0.34 (Cr203	0.00	0.00	0.00	0.00	FeO	0.28	0.04	0,00	0.25	0.02
FeO	12.5	16.50	14.88	20.9	12,00	15.7	9.3	6.92	7.38	16.78	12.49	13.36 I	Fe2O3	5.32	6.55	13.20	7.07	MnO	0.00	0.13	0.06	0.00	60.0
MgO	14.74	10.23	11.4	5.05	16.83	12.47	17.18	19.71	17.59	7.64	13.2	13.59	MnO	0.03	0.00	0.14	0.00	MgO	0.17	0.01	0.00	0.00	0.19
CaO	8.58	8.72	7.98	9.7	10.08	6.66	12.97	11.98	11.47	11.53	7.97	11.98	MgO	0.05	0.21	0.13	0.26	CaO	6.32	0.06	4.6	3.84	6.29
Na2O	3.57	4.18	2.81	4.56	2.39	4.99	1.13	1.59	1.73	3.06	3.65	1.04 0	CaO	24.27	23.78	23.11	23.98	Na2O	8.76	12.58	8.6	10.28	8.94
K20	0.03	0.07	0.08	0,00	0.22	0,00	0,00	0.07	0,00	0.4	0.16	0.15 7	Total	97.91	97.30	<i>PT.77</i>	98.37	K20	0.07	0.03	0.05	0.10	0.03
Total	97.88	97.65	96.30	96.87	98.35	99.13	96.45	98.09	97.17	97.75	96.59	96.41						Total	100.87	99.82	100.79	99.57	100.23
												-1	Si	5.996	5.977	5.993	5.940						
Si	7.807	7.110	7.287	5.969	7.748	7.192	7.793	7.729	7.490	5.986	7.240	7.025 7	Ti	0.000	0.022	0.013	0.015	Si	2.679	2.984	2.782	2.793	2.678
AI(IV)	0.193	0.890	0.713	2.031	0.252	0.808	0.207	0.271	0.510	2.014	0.760	0.975	AI	5.365	5.213	4.411	5.212	Ti	0.000	0.000	0.002	0.003	0.003
sumT	8.000	8.000	8.000	8.000	8.000	8.000	8.000	8.000	8.000	8.000	8.000	8.000	Cr	0.000	0.000	0.000	0.000	AI	1.295	066.0	1.231	1.178	1.290
AI(VI)	0.355	0.709	0.703	1.136	0.045	0.508	0.152	060.0	0.417	1.105	0.645	0.332 1	Fe3	0.619	0.770	1.583	0.824	Fe2	0.010	0.001	0.000	0.009	0.001
Ti(VI)	0.015	0.010	0.017	0.003	0.005	0.017	0.001	0.001	0.011	0.064	0.000	0.032	Mn	0.003	0.000	0.020	0.000	Mn	0.000	0.005	0.002	0.000	0.003
Fe3+	0.203	0.215	0.702	0.346	0.458	0.896	0.000	0.164	0.141	0.112	0.622	0.503 1	Mg	0.011	0.050	0.030	0.059	Mg	0.011	0.001	0.000	0.000	0.012
Mg	3.125	2.237	2.455	1.155	3.536	2.616	3.715	4.091	3.691	1.707	2.822	2.961	Ca	4.019	3.980	3.947	3.977	Ca	0.300	0.003	0.216	0.183	0.300
Fe2+	1.284	1.810	1.096	2.335	0.956	0.951	1.128	0.642	0.728	1.991	0.875	1.130	Total	15.380	15.170	15.934	15.129	Na	0.752	1.072	0.730	0.889	0.772
Mn	0.018	0.019	0.027	0.025	0.000	0.012	0.003	0.012	0.012	0.021	0.036	0.042						К	0.004	0.002	0.003	0.006	0.002
sumC	5.000	5.000	5.000	5.000	5.000	5.000	4.999	5.000	5.000	5.000	5.000	5.000						Total	5.051	5.058	4.966	5.062	5.062
Ca	1.307	1.371	1.235	1.594	1.521	1.004	2.000	1.787	1.730	1.851	1.224	1.876											
Na	0.693	0.629	0.765	0.405	0.479	0.996	0.000	0.213	0.270	0.149	0.776	0.124						Anorthite	27.84	0.24	22.70	16.86	27.53
sumB	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000	2.000						Albite	69.82	98.94	76.78	81.76	70.79
Na	0.291	0.560	0.022	0.951	0.174	0.365	0.318	0.216	0.202	0.740	0.239	0.171						Orthoclas	0.35	0.14	0.31	0.52	0.18
К	0.005	0.013	0.015	0.000	0.040	0.000	0.000	0.013	0.000	0.077	0.029	0.028											
sumA	0.296	0.573	0.037	0.951	0.214	0.365	0.334	0.229	0.202	0.817	0.268	0.199											
catsum	15.296	15.573	15.037	15.951	15.214	15.365	15.333	15.229	15.202	15.817	15.268	15.199											
	WIN	Mgkat	BarA	Bar Al-Feparg	Act	Febar	Act	Act	Mghor Al-Feparg	Feparg	Bar	Mghor											

Amph bd: microboudin in amphibolite; amph lw: amphibolite with pseudomorphs after lawsonite; amph ep: epidote amphibolite; met gb; metagabbro. Amphibole structural formula recalculated using the method of Richard and Clark (1990), based on 23 oxygens. Total number of cations = 15 excluding K. Nomenclature after Leake et al. (1997). Unit formula of epidote based on 25 oxygens, and of feldspar on 8 oxygens.

Table 2 - Selected SEM-EDS analyses from eclogite samples of the Antrona ophiolite.

		AMPHIBOLE						CLINOPYRO	XENE		
SAMPLE	C1	C1	C1	C1	C1	SAMPLE	C1	C1	C1	C1	C14
Analysis	C1AN3	C1CAN1	C1CAN2	C1CAN3A	C1CAN4	Analysis	C1APX2*	C1APX4*	C1EPX1*	C1EPX2*	C14BPX1
Occurrence	incl. in gar	rim/gar	core	core	rim/gar	Occurrence	rim	core	rim	rim	rim
SiO2 wt%	57.61	56.12	56.22	54.57	54.34	SiO2 wt%	55.05	54.82	54.76	54.71	56.26
TiO2	0.01	0.01	0.01	0.00	0.00	TiO2	0.03	0.03	0.02	0.06	0.07
Al2O3	7.81	3.26	2.71	4.85	3.81	Al2O3	4.53	2.98	3.96	4.18	7.20
Cr2O3	0.00	0.00	0.01	0.00	0.00	Cr2O3	0.00	0.03	0.00	0.06	0.00
FeOt	14.72	11.64	12.61	13.20	14.07	FeOt	10.18	10.74	10.94	10.41	6.76
MnO	0.00	0.00	0.01	0.01	0.01	MnO	0.00	0.07	0.02	0.04	0.02
MgO	10.18	15.59	15.97	14.54	14.82	MgO	9.32	10.09	9.54	9.84	9.22
CaO	1.47	8.00	8.64	8.47	8.70	CaO	15.70	16.79	15.27	16.14	14.55
Na2O	6.19	2.60	2.07	2.57	2.23	Na2O	5.67	4.55	5.38	5.05	6.43
K2O	0.00	0.01	0.01	0.00	0.00	K2O	0.00	0.00	0.00	0.00	0.00
Tot	97.99	97.23	98.26	98.21	97.98	Tot	100.48	100.09	99.88	100.48	100.51
Si	8.010	7.900	7.877	7.645	7.665	Si	1.990	2.005	1.998	1.986	2.003
Al	0.000	0.100	0.123	0.355	0.335	Al4	0.010	0.000	0.002	0.014	0.000
SumT	8.010	8.000	8.000	8.000	8.000	tet	2.000	2.005	2.000	2.000	2.003
Al	1.281	0.441	0.325	0.447	0.299	Al6	0.183	0.129	0.169	0.164	0.302
Cr	0.000	0.000	0.001	0.000	0.000	Fe2	0.086	0.137	0.121	0.115	0.063
Fe3	0.367	0.364	0.356	0.606	0.646	Fe3	0.222	0.192	0.212	0.200	0.138
Ti	0.001	0.001	0.001	0.000	0.000	Mg	0.502	0.550	0.519	0.533	0.490
Mg	2.109	3.271	3.335	3.036	3.115	Mn	0.000	0.002	0.000	0.001	0.001
Fe2	1.242	0.923	0.983	0.911	0.940	Ti	0.001	0.001	0.000	0.002	0.002
Mn	0.000	0.000	0.000	0.000	0.000	Cr	0.000	0.001	0.000	0.002	0.000
Ca	0.000	0.000	0.000	0.000	0.000	Oct	0.994	1.011	1.023	1.017	0.995
SumC	5.000	5.000	5.000	5.000	5.000	Xoct	0.000	0.011	0.023	0.017	0.000
Mg	0.000	0.000	0.000	0.000	0.000	Ca	0.608	0.658	0.597	0.628	0.555
Fe2	0.103	0.084	0.139	0.029	0.074	Na	0.398	0.323	0.381	0.355	0.444
Mn	0.000	0.000	0.001	0.001	0.001	M2	1.006	0.991	1.000	1.000	0.998
Ca	0.219	1.207	1.297	1.271	1.315						
Na	1.669	0.710	0.562	0.698	0.610	Jd	18.3	12.9	16.9	15.9	30.0
SumB	1.990	2.000	2.000	2.000	2.000	Ac	22.2	19.2	21.2	19.5	13.8
Ca	0.000	0.000	0.000	0.000	0.000	Wo	30.3	33.2	29.9	31.8	28.1
Na	0.000	0.000	0.000	0.000	0.000	En	25.0	27.8	26	27.0	24.7
К	0.000	0.002	0.002	0.000	0.000	Fs	4.3	6.9	6.1	5.8	3.2
SumA	0.000	0.002	0.002	0.000	0.000						
tot cat	15.000	15.002	15.002		15.000						
	121300	15.502	10.002	12.500	10.000						

Table 3 - Selected electron microprobe analyses of amphibole and clinopyroxene from eclogite samples of the Antrona ophiolite.

15-K Glaucophane Winchite Actinolite Winchite Actinolite

Amphibole structural formula recalculated using the method of Richard and Clark (1990), based on 23 oxygens. Total number of cations = 15 excluding K. Nomenclature after Leake et al. (1997). Clinopyroxene formula based on 6 oxygens.

tions at the core as well as at the rim of crystals (Table 3). Amphibole inclusions in garnet are glaucophane (Table 3).

Clinopyroxene from eclogite shows compositions ranging from omphacite to aegirine-augite (Table 3), according to Morimoto's (1989) classification.

Garnet in eclogite is characterized by chemical zoning with grossular-rich cores and pyrope-rich rims (Table 4).

P-T conditions of metamorphism

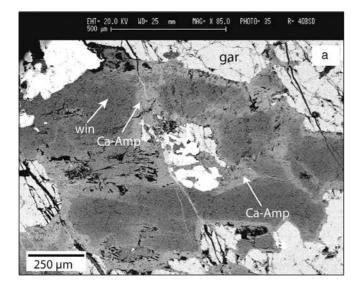
Temperature and pressure conditions of the Alpine metamorphic peak were estimated on amphibole-bearing eclogite samples from the study area. P-T values are constrained by the garnet-Na-clinopyroxene equilibria and the jadeite content in Na-clinopyroxene. Temperature was evaluated as the average of the four more recent calibrations for the garnet-clinopyroxene thermometer (Powell, 1985; Ai, 1994; Krogh, 1998; Ravna, 2000). Six garnet-clinopyroxene pairs were analised and temperatures of T = $372 \pm 50^{\circ}$ C for a nominal pressure of P = 1 GPa and T =

 $386 \pm 50^{\circ}$ C for a nominal pressure of P = 1.5 Gpa were obtained. We found pyroxenes with Jd₃₀ as the maximum jadeite content (see Table 3) suggesting P > 1 GPa (Holland, 1983).

DISCUSSION

Rock assemblage and lithostratigraphy

The ophiolite suite exposed in the Antrona Valley consists of serpentinized (mantle) peridotite, metagabbro, metabasalt and metasedimentary cover, largely representing the typical "Penrose"-type ophiolite (Anonymous, 1972). On the basis of field data and petrography, a new recognition of the ophiolite rock types was performed, in order to define the stratigraphy of this portion of oceanic lithosphere. More detailed informations would be obtained by integrating the present results with bulk rock analyses (Tartarotti et al., in prep.). Metabasaltic rocks, defined so



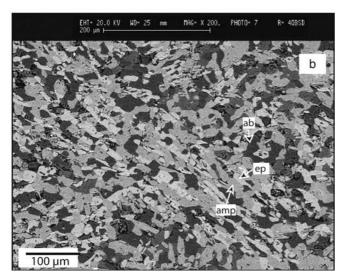


Fig. 6 - BSE images of amphibolite and eclogite samples. a) Chemical zoning in eclogite amphibole: Na-Ca amphibole (win = winchite) corresponds to darker core; lighter Ca-amphibole (Ca-Amp) forms the rims; gar = garnet. b) Detail of the "sheaf"-like structure observed in light green boudins of the microboudin-bearing amphibolites. amp= amphibole; ep = epidote; ab = albite.

far simply as "amphibolites" are tentatively subdivided into different types, on the basis of their mineral composition and microstructural features. New findings are: i) relics of magmatic structures in some amphibolites, inferred to be pillow lavas or pillow breccia; ii) lawsonite pseudomorphs-bearing amphibolites; iii) that epidote-amphibolite of the Antrona Valley derived from lawsonitebearing amphibolite.

Metamorphic evolution

Petrographic observations and mineral analyses give information about the Alpine tectonometamorphic evolution for the studied ophiolite rocks, summarized in Fig. 9. By integrating microstructural features and structures observed at the meso-scale, at least four Alpine ductile deformation phases (D1-D4) were recognized. These phases produced four metamorphic foliations (S1-S4). Four stages of mineral crystallization (B1-B4) were recognized.

D1 is related to relict microstructures older than D2 (re-

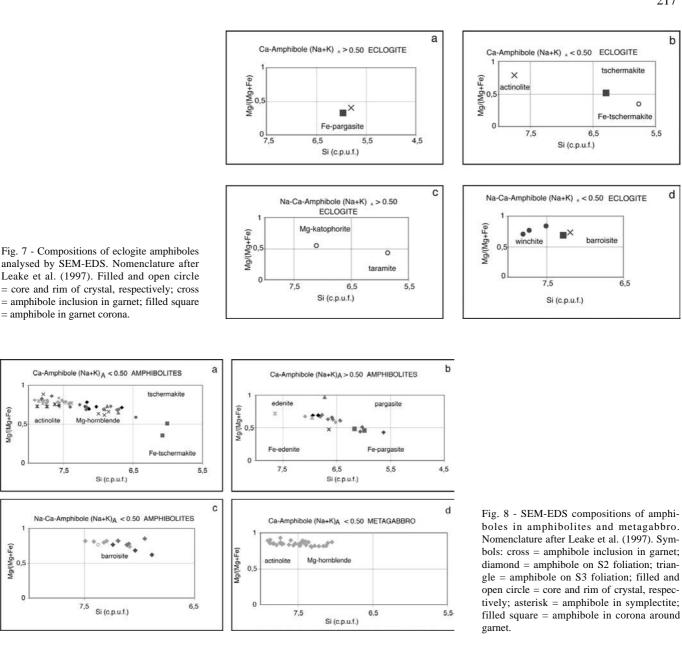
sponsible for the main foliation S2). Such microstructures are represented by glaucophane inclusions (B1) inside garnet in the amphibole-eclogite samples. Glaucophane inclusions, however, are scattered and do not define an internal foliation within garnet. Internal foliation marked by rutile and quartz was instead observed inside garnet poikiloblasts in micaceous metaquartzites (Turco, 2004). We infer that glaucophane inclusions and rutile + quartz foliation represent relict of the prograde Alpine metamorphic path likely occurred under blueschist facies conditions. D2 is responsible for the main regional foliation S2 recognizable in the field, mostly defined by Ca-Na- and Ca-amphibole. Before the onset of D2, step B2 (eo-Alpine stage) is responsible for the crystallization of high pressure mineral assemblages including Na-clinopyroxene + winchite + garnet + rutile in mafic rocks (Fig. 9). Step B3 (meso-Alpine stage) produced Ca-Na amphiboles at winchite rims in eclogite and Ca-amphibole + epidote + biotite + plagioclase in amphibolites, marking the S2 foliation. B4 (neo-Alpine stage) is characterized by the growth of actinolite + chlorite + epidote + albite defining the S3 and S4 foliations under greenschist facies conditions. In some samples, chlorite and epidote are not associated with a foliation, when replacing garnet. S3 and S4 foliations are not pervasive and are defined as crenulation cleavage or S-C structures. Finally, B4 was followed by growth of prehnite filling veins cutting through all previous structures.

Microstructural evidence regarding the first appearance of lawsonite in the studied rocks is scarce: relics of (pseudomorphed) lawsonite inside garnet were not observed. Evidence of Na-clinopyroxene-lawsonite coexistence is lacking (Na-clinopyroxene was found in eclogites and not in amphibolites bearing pseudomorphs after lawsonite). Indeed, bulk rock chemistry, in addition to pressure, temperature and volatile components in the system, are among the main factors controlling lawsonite formation (e.g., Pognante, 1989 and refs. therein; Bucher et al., 2005). Such scarce textural evidence prevents a correct definition of the relative chronology of deformation vs. metamorphism. On the basis of textural relations, we infer that the growth of lawsonite porphyroblasts in amphibolites took place before or during D2 which produced S2 foliation defined by (B3) Na-Ca and Ca-amphibole. Consequently, lawsonite (± rutile; see Fig. 5d) is inferred to grow before the onset of the amphibolite or epidote-amphibolite facies conditions.

A qualitative P-T diagram showing the metamorphic evolution the Antrona ophiolite rocks as deduced from the stability conditions of the observed mineral parageneses is illustrated in Fig. 10. The peak P-T condition as estimated in the eclogite samples is also reported (see "B2" in Fig. 10). The prograde event (B1) likely occurred under blueschist facies (as attested by glaucophane inclusions in garnet from eclogite samples) followed by high pressure (eclogitic) metamorphic peak, attested by the omphacite-garnet pair (B2). A possible coexistence of the omphacite-lawsonite pair under eclogite facies conditions cannot be ruled out, if we consider the P-T values obtained by thermobarometric calculation in the eclogite samples. In fact our results (T = $372 \pm 50^{\circ}$ C for $P \ge 1$ Gpa) are compatible with the stability field of lawsonite (low-T side of the reactions 1 and 2 in Fig. 10). Previous works on eclogites in the studied area suggested that metabasic rocks were equilibrated under P = 1.4 - 1.6 GPa and $T = 480-720^{\circ}C$ (Colombi and Pfeiffer, 1986). These results have been obtained on garnet - omphacite pairs, follow-

b

d



ing the Ellis and Green (1979)'s thermometric calibration. Nevertheless, the suggested temperature span appears to be unreasonably large. In addition, no microchemical analyses or other arguments have been provided by Colombi and Pfeiffer (1986) to explain such a large temperature span. Many analyses (e.g. Lardeaux et al., 1986) suggested that the temperature span obtained with this method may considerably be reduced with a careful textural selection of the analysed sites. Moreover, the Ellis and Green (1979)'s calibration for the garnet-clinopyroxene thermometer probably overstimates temperature (e.g. Green and Adams, 1991) and thus it was not utilised in the present study.

According to Pfeifer et al. (1989), the peak metamorphism of the Antrona metabasic rocks was followed by amphibolite facies conditions, estimated at $T = 550^{\circ}C$ in metasediments, and then by greenschist facies conditions. According to Carrupt and Schlup (1998), the retrogressive greenschist stage was followed by an increase in temperature, inferred from the compositional zoning of amphibole and plagioclase in mafic rocks. In the present study, the retrograde metamorphic evolution may be inferred from the incipient breakdown of high pressure minerals in the eclogite samples and from the amphibolites samples, that are characterized by epidote-amphibolite/amphibolite facies mineral assemblages (step B3 in Figs. 9 and 10). These mineral assemblages were overprinted by greenschist facies minerals (B4), followed by prehnite-pumpellyite facies mineral assemblages (prehnite-filled veins). Our petrographic observations and mineral chemistry data are consistent with Pfeifer et al. (1989)'s estimates for the retrogressive evolution of the Antrona ophiolite.

CONCLUDING REMARKS AND REGIONAL IMPLICATIONS

Field data and petrographic observations allow a new definition of the ophiolite rock types exposed in the Antrona Valley. New findings are:

- relics of magmatic structures in some amphibolites, inferred to be pillow lavas or pillow breccia;
- lawsonite pseudomorphs-bearing amphibolites;
- ٠ the epidote-amphibolites of the Antrona Valley derived from lawsonite-bearing amphibolite;

SAMPLE	C1	C1	C1	C1	C1	C1	C1
Analysis	C1CGT1	C1CGT3	C1CGT4	C1AGT2*	C1AGT3*	C1EGT1*	C1EGT2*
Occurrence	rim	int rim	core	rim	rim	rim/cpx	int rim/cpx
SiO2 wt%	38.56	38.17	37.96	38.39	38.09	38.40	38.08
TiO2	0.09	0.06	0.14	0.07	0.11	0.05	0.09
Al2O3	20.51	20.61	20.46	20.89	20.44	20.43	20.60
Cr2O3	0.00	0.00	0.00	0.04	0.02	0.00	0.00
FeOt	31.74	30.82	28.96	32.12	30.67	31.54	30.86
MnO	1.21	0.23	1.47	0.13	1.28	0.47	0.16
MgO	3.82	2.02	1.47	2.46	1.28	2.73	2.13
CaO	5.95	9.29	11.17	7.87	10.29	7.40	8.83
Na2O	0.00	0.00	0.00	0.01	0.06	0.07	0.00
K2O	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Tot	101.88	101.21	100.43	101.97	101.23	101.10	100.76
Si	3.008	3.007	3.008	3.004	3.008	3.022	3.011
Ti	0.005	0.004	0.008	0.004	0.007	0.003	0.005
Al	1.886	1.914	1.911	1.927	1.902	1.895	1.920
Cr	0.000	0.000	0.000	0.003	0.001	0.000	0.000
Fe3	0.109	0.083	0.081	0.067	0.090	0.102	0.075
Fe2	1.961	1.947	1.839	2.035	1.936	1.974	1.966
Mn	0.080	0.016	0.018	0.009	0.019	0.031	0.011
Mg	0.444	0.238	0.174	0.286	0.151	0.321	0.251
Ca	0.497	0.784	0.949	0.660	0.871	0.624	0.748
Na	0.000	0.000	0.000	0.001	0.009	0.011	0.000
Tot	7.990	7.991	7.987	7.995	7.993	7.982	7.987
Pyrope	14.89	7.96	5.83	9.58	5.06	10.87	8.44
Almandine	65.76	65.25	61.73	68.07	65.05	66.91	66.05
Grossular	11.18	22.12	27.79	18.59	24.67	15.99	21.37
Spessartine	2.68	0.52	0.59	0.29	0.63	1.06	0.36
Andradite	5.48	4.15	4.06	3.34	4.52	5.17	3.77

Table 4 - Selected electron microprobe analyses of garnet from eclogite samples of the Antrona ophiolite.

* utilised for thermobarometric calculations. Structural formula calculated on the basis of 12 oxygens.

• the P-T evolution is characterized by a prograde blueschist-facies stage and by an eclogitic peak occurred at $T = 372 \pm 50$ °C and $P \ge 1$ GPa. This P-T condition is consistent with the stability of lawsonite, that was observed as pseudomporphs in amphibolites, but not in omphacite-garnet-bearing rocks. The retrograde evolution of the Antrona metabasites occurred under epidote-amphibolite/amphibolite facies overprinted by greenschist facies conditions. Late minerals associated with brittle deformation grew under prehnite-pumpellyite facies conditions.

The inferred metamorphic evolution for the Antrona Unit may be compared, at least in part, to that suggested for the Zermatt-Saas metaophiolite (e.g., Ernst and Dal Piaz, 1978; Barnicoat and Fry, 1986; Fry and Barnicoat, 1987; Bucher et al., 2005): both units exhibit prograde Alpine blueschist mineral assemblages followed by an eclogitic peak, typical for subduction-related metamorphism. However, the P-T condition of the eclogitic peak in the two units are not easily comparable, due to the large span of P-T estimates obtained so far for the Zermatt-Saas eclogites, ranging from 450-780°C and from 1-2.2 GPa (e.g., Ernst and Dal Piaz, 1978; Oberhänsli, 1980; Barnicoat and Fry, 1986; Fry and Barnicoat, 1987; Cartwright and Barnicoat, 2002).

The retrograde path of the Antrona ophiolite differs from those proposed by Barnicoat and Fry (1986) and Fry and Barnicoat (1987) for the Zermatt-Saas Unit. This latter is characterized by an initial rapid cooling to the lawsoniteeclogite field, before the onset of blueschist (glaucophane growth) and then greenschist facies conditions. Although not well contrained in its initial part, decompression path of the Antrona ophiolite is dominated by epidoteamphibolite/amphibolite facies mineral assemblages. Consequently, even if a common prograde path may be envisaged for the Zermatt-Saas and Antrona ophiolite, their respective exhumation histories probably became different, accounting for their different tectonic positions in the Alpine nappe pile.

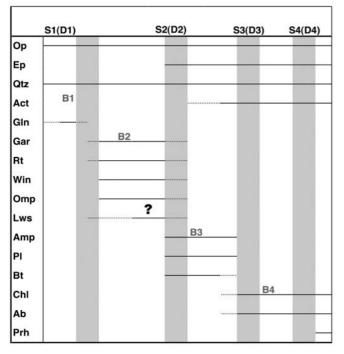


Fig. 9 - Deformation (D) and foliation (S) vs. mineral growth (B) inferred for the studied ophiolite samples from the Antrona ophiolite. Mineral abbreviations according to Kretz (1983).

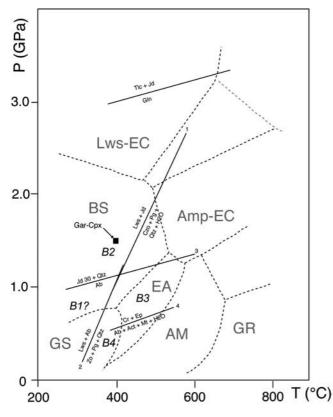


Fig. 10 - Petrogenetic grid with critical reactions defining the metamorphic evolution of the Antrona ophiolite rocks. B1 – B4 represent stages of mineral crystallization as defined in figure 9. Black square indicates the P-T conditions estimated as the average of four gar-cpx thermometers (Powell, 1985; Ai, 1994; Krogh, 1998, and Ravna, 2000), and by considering the maximum jadeite content in clinopyroxene (Holland, 1983). Metamorphic facies fields after Spear (1993). Reaction curves: 1) lws+jd = zo+pg+qtz+H2O (Heinrich and Althaus, 1980); 2) lws+ab = zo+pg+qtz+H2O (Heinrich and Althaus, 1980); 3) omp (jd30) + qtz = ab (Holland, 1983) 4) crossite+ep = ab+act+mag+H2O (Maruyama et al., 1986). Mineral abbreviations according to Kretz (1983).

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