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(Article begins on next page)

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Asbestiform sepiolite coated by aliphatic hydrocarbons from Perletoa, Aosta Valley Region (Western Alps, Italy): characterization, genesis and possible hazards

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Abstract

An atypical asbestiform sepiolite occurrence with exceptionally long fibres wrapped by a sheath of aliphatic hydrocarbons was found in the Gressoney Valley (Italian Western Alps) while monitoring asbestos presence in outcropping serpentinite rocks.

Microscopic and FT-IR analyses proved that these fibres, apparently up to several cm long, are formed by bundles of thinner fibrils (average length: 150 μm) potentially dispersible in the environment. When observed with TEM these fibrils show a rhomboidal to parallelogram cross section ($< 1 \mu\text{m}$), whose surfaces are mostly covered by an aliphatic hydrocarbons film – an association never reported in literature. The sepiolite fibrils and their organic coating probably originated in sequential steps from precipitation of Si/Mg rich hydrothermal fluids, resulting from serpentinization of olivine and clinopyroxene, and Fischer-Tropsch-type reaction. Presence of hydrocarbons implies serious consequences on the sepiolite habit, as the organic wrap interacts with the fibrils surface reducing the amount of adsorbed water and favouring fragmentation of thicker units into thinner ones, due to an ‘opening’ process implying separation along z and cleavage on (110). This defibrillation mechanism, coupled to the extraordinary length, further increases these fibrils aspect ratio (length/width $\gg 3$) thus amplifying their potential danger for human health when air dispersed and breathed.

Keywords: sepiolite, aliphatic hydrocarbons, Fischer-Tropsch-type reaction, hydrothermal system, noxiousness hazard.

1. Introduction

Sepiolite is a fibrous, trioctahedral phyllosilicate clay mineral belonging to the palygorskite-sepiolite group. Its natural deposits occur as chemical sediments or reconstituted former sedimentary clays in epicontinental and inland seas and lakes, where somewhat saline groundwater discharges under semi-arid climatic conditions (Eugster and Hardie, 1975; Mayayo *et al.*, 1998). Other sepiolite occurrences form by hydrothermal alteration of basaltic glass, volcanoclastic sediments or previous clays in the open ocean (in association with fore-arc basins and ocean ridges) or in calcareous soils by direct crystallization (Galán and Pozo, 2011). An authigenic clay mineral, sepiolite can form *in situ* through direct precipitation from solutions containing dissolved ionic species (namely Si and Mg; Trauth, 1977; Jones and Galan, 1988; Weaver, 1984) either in lacustrine (Chahi *et al.*, 1997) or perimarine (Velde, 1985) environments in conditions of high alkalinity (pH: 8 - 9.5) and medium salinity. Direct precipitation from low-temperature hydrothermal solutions in continental environments were described (Imai and Otsuka, 1984) as well as precipitation, on or below the sea floor, from Mg/Si enriched low T (< 100°C) solutions formed by the interaction of ultramafic rocks with seawater circulating in the crust, particularly in transform zones (Bonatti *et al.*, 1983). Other peculiar findings included possible biogenic origin induced by microorganisms (Leguey *et al.*, 2010; Cuevas *et al.*, 2012). Although not a common mineral, sepiolite deposits can be found worldwide – especially in Spain (Neogene

lacustrine sequences of the Madrid basin; Brell *et al.*, 1985; Ordoñez *et al.*, 1991) and Turkey (continental lacustrine deposits of the Ezkisehir Basin; Ece and Çoban, 1994), but also USA (Amargosa Desert deposit), Kenia and Tanzania (Amboseli deposit), China (Guanshan deposit), Greece (Ventzia basin deposit) and Somalia (El Bur deposit) (Galán and Pozo, 2011).

Sepiolite has various and useful industrial applications such as lubricant, catalyst, adsorbent, cleaning compress for restoration and cat litter (Álvarez *et al.*, 2011; Lòpez-Galindo *et al.*, 2011). In addition sepiolite may be used as an asbestos substitute in building and seal materials, friction compounds for automotive brakes and high temperature insulators (Noda *et al.*, 2009; Solebello, 2009) due to its habit and modest fibre length. Almost all commercially sold specimens have fibres < 5 µm long – the threshold of non-carcinogenicity accepted by the International Agency for Research on Cancer, World Health Organization (1997) – sometimes as the result of preliminary grinding. However, sepiolite occurrences from all over the world show high variability in cristallinity, fibre length and texture (Suárez and García-Romero, 2012; García-Romero and Suárez, 2013); samples with unusually long fibres (> 10 µm) were seldom identified and signalled for their carcinogenic potential (Pott *et al.*, 1990; Pott *et al.*, 1991; Bellman *et al.*, 1997).

The structure of sepiolite [crystal-chemical formula $Mg_8Si_{12}O_{30}(OH)_4(OH_2)_4 \cdot 8H_2O$, s.g. Pbnm], first discussed by Preisinger (1959, 1963) and Brindley (1959), show a continuous waving tetrahedral sheet (T) alternated (along X) to a discontinuous octahedral one (O) forming ribbons elongated in the Z-axis direction. Such an arrangement, described as a framework of chessboard connected TOT ribbons (Ferraris *et al.*, 2008), causes the structure to be crossed by Z-elongated tunnels (10.6 x 3.7 Å)

filled by weakly bound zeolitic H₂O and exchangeable ions. Tightly bound structural OH₂ (Bailey *et al.*, 1980; Guggenheim and Krekeler, 2011) completes the coordination of Mg ions at the borders of the O ribbons, The structure of sepiolite differs from that of palygorskite, although recently intermediate forms were described which suggest the possible existence of a continuous compositional variation between the end-members of the group (Garcia-Romero *et al.*, 2007; Garcia-Romero and Suarez, 2010).

This study describes an atypical occurrence of sepiolite from Perletoa village, in the Gressoney Valley near the town of Aosta (Western Alps, Italy), showing unusually long fibres wrapped by a sheath of aliphatic hydrocarbons. The finding of asbestiform sepiolite in NW Italian outcropping rocks is not a novelty. Long fibre samples were detected in specific areas of the Valle d'Aosta, Piemonte and Liguria Regions (Belluso and Sandrone, 1989; Belluso *et al.*, 1995). The genesis of organic matter (OM)-bearing clays (i.e. black shales or bituminous rocks) was reported from different sedimentary settings (Emeis and Weissert, 2009) or by weathering of mafic to ultramafic lithologies (Dos Anjos *et al.*, 2010). Besides, hydrocarbon bearing clays were discovered in some metasomatized oxide-rich gabbros (Ciliberto *et al.*, 2009) or in a diapiric intrusion in a diatremic tuff-breccia deposit, precipitated from a mixture of hot Si-rich hydrothermal fluids (350-400° C) and cold seawater (Manuella *et al.*, 2012). Though sepiolite traces were detected in the latter study, no mention of its coupling to aliphatic hydrocarbons has ever been made so far. This work focuses therefore on such a peculiar association and consequent interaction, which affects the clay mineral habit possibly increasing its potential noxiousness for human health in case its fibres are airborne and breathed in large amounts.

2. Experimental

1.1 Materials and geological setting

The site of collection of the investigated sepiolite is located near Perletoa village, in the Gressoney Valley near the town of Aosta (Aosta Valley Region, Western Alps, Italy; 45° 48' 5" N, 7° 49' 25" E) (Fig. 1). This clay mineral was noted during systematic sampling to evaluate presence of asbestos and asbestiform minerals in massive to foliated serpentinite rocks from the main ultramafic bodies of the Italian Western Alps (Belluso *et al.*, 1995). The analyzed sepiolite specimen was collected in the area of the Combin Zone (Geological Map of Aosta Valley, available at http://geonavsct.partout.it/pub/GeoNavSCT/index.html?repertorio=carta_geologica_100k), from a small serpentinitic lens outcropping (as peridotite, prasinite, amphibolite and eclogite lenses) in dominant calcschists.

(INSERT FIGURE 1)

2.2 Methods

Scanning electron microscopy (SEM) observations were obtained with a FEG SEM JEOL JSM6320F instrument and a SEM Stereoscan 360, Cambridge Instrument. The samples, coated by a 50 Å thick Au layer to allow conductivity, were attached to step brass stubs to allow examination of the fibres. Chemical characterization was performed by electron probe microanalysis (EPMA) using an EDS Link Pentafet, Oxford instrument. Due to the difficulties in analyzing fibrous samples, analyses were collected

on 10 x 10 μm^2 areas on a pressed, sintered and carbon-coated sepiolite pellet obtained with a press designed to prepare samples for infrared spectroscopy (operating conditions: 50 s counting time, 15 kV accelerating voltage, 25 mm working distance, 300 pA beam current). Collected data were processed with the Microanalysis Suite Issue 12, INCA Suite Version 4.01 and calibrated on natural mineral standards using the ZAF correction method. The weight% sums of oxides, showing no relevant heterogeneity, were averaged to obtain a reliable crystal-chemical formula.

A 120 kV transmission electron microscopy (TEM Philips CM12 working at 120 kV, LaB₆ filament, double tilt holder, equipped with an energy dispersive spectrometer EDAX Genesis 2000 System, TEM Quant Software PV8206/31 procession system) was used to examine crushed and ion thinned samples. Only medium to high TEM magnification can be performed at higher electron energy (lower ionisation), owing to the high instability of the fibrous sepiolite specimen under the electron beam. A TEM JEOL 2000FX (200 kV accelerating voltage, side entry, double-tilt $\pm 30^\circ$, point-to-point resolution of 2.8 Å), equipped with a low-light camera (LHESA EM LH4086) with a YAG converter, was used. TEM specimens, cut perpendicular and parallel to the elongation of the fibres, were extracted from standard petrographic, uncovered thin sections (30-40 μm thick) embedded with epoxy resin and attached by Lakeside resin onto a glass slide. The microdrilled disks (attached to single-hole copper TEM slots) were thinned by ion beam milling (GATAN 600 Duomill working with Ar at room temperature, 5 kV and 0.50 mA per gun, 15° and finish angle of 12°). A powdered sample was ground in an agate and pestle mortar with isopropyl alcohol and then sonicated; two drops of the resulting suspension were deposited on a copper mesh grid previously coated with a 200 Å carbon film. At medium to high magnification, several

morphological and structural pictures were obtained. Selected area electron diffraction (SAED) were displayed but not photographed owing to very rapid fibres amorphization.

Thermogravimetric data were collected on a 15 mg sample with a simultaneous TGA/DSC SDT Q600, TA Instruments, in air flow with a heating rate of $10\text{ }^{\circ}\text{C min}^{-1}$ from room temperature to $900\text{ }^{\circ}\text{C}$. DSC heat flow data were dynamically normalized using the instantaneous sample weight at any given temperature.

For infrared (IR) spectroscopy, the sepiolite was preliminarily activated in vacuum (pressure below 5×10^{-4} mbar) at room temperature, $120\text{ }^{\circ}\text{C}$ and $150\text{ }^{\circ}\text{C}$ for 1.5 hours each to progressively (and reversibly) remove the zeolitic H_2O . The resulting powder was used to make a pellet that was inserted in an IR cell and further heated in vacuum (pressure $< 5 \times 10^{-4}$ mbar) at $120\text{ }^{\circ}\text{C}$ and $150\text{ }^{\circ}\text{C}$. IR absorption spectra were collected on a FTIR Bruker Vector 70, with a resolution of 2 cm^{-1} and collecting 64 scans for each spectrum. Data were collected at the different investigated T steps (room T , $120\text{ }^{\circ}\text{C}$ and $150\text{ }^{\circ}\text{C}$) under controlled atmosphere.

For synchrotron X-ray powder diffraction (XRPD) and Rietveld refinement, the sepiolite specimen was initially purified by mechanically removing incidental macroscopic contaminants (serpentine, calcite and incidental pollutants) under the binocular stereo-microscope. The resulting fraction was dispersed in deionized water, thus possibly isolating the lighter suspended fraction from the heavier impurities [i.e. quartz and/or calcite] (Giustetto and Chiari, 2004). Massive purified samples were hand-ground in an agate mortar and crushed to powder. Conventional XRPD showed sepiolite to be the only detectable phase.

Synchrotron XRPD data were collected at room T at the European Synchrotron Radiation Facility (ESRF) in Gr noble (France), on the GILDA (General-purpose

Italian beam-Line for X-ray Diffraction and Absorption) beamline, using a wavelength of $\lambda = 0.6530 \text{ \AA}$ and Debye-Scherrer geometry. The specimen was loaded into a 1 mm diameter quartz-glass capillary to avoid preferred orientation. Collection time was 120 sec. Diffracted beams were collected with an area detector (MAR345) accessing d -values from 0.8 to 80 \AA .

The GSAS software package (Larson and Von Dreele, 2007) and the EXPGUI graphical user interface (Toby, 2001) were used for Rietveld refinement. Background was fitted using a 14-term shifted-Chebyshev function and peak profiles modelled with a pseudo-Voigt function as parameterized by Thompson *et al.* (1987), with asymmetry corrections according to Finger *et al.* (1994). Initially, only the background, scale factor, zero and unit-cell parameters were refined. Later, fractional coordinates and occupancy factors for all atoms were refined, at first in alternate cycles (to minimize correlations) and then simultaneously. Soft constraints were imposed on tetrahedral and octahedral bond lengths and angles and progressively reduced. Isotropic displacement parameters (Uiso) for all atoms were adjusted using overall constraints for each chemical species. Preferred orientation for the (110) reflection (attenuated by data collection in capillary) was treated with the March-Dollase model (March, 1932; Dollase, 1986). Consistency of the refined structure was constantly checked through screening of the GOF (Goodness-Of-Fit) parameters, graphical fit between observed and calculated profiles and graphical representation of the refined model (Moldraw: Ugliengo *et al.*, 1993). The lower soft-constraint weighting factor yielding reasonable bonds in tetrahedrons ($F = 10$) was kept to avoid unrealistic bond distances and polyhedrons distortions (Post *et al.*, 2007; Post and Heaney, 2008; Giustetto *et al.*, 2011a; Giustetto and Compagnoni, 2011).

3. Results and discussion

3.1 Macroscopic and stereomicroscopic observation

Even at the macroscopic scale, the asbestiform habit of the Perletoa sepiolite is evident. Long and apparently thick fibres (from 5 mm up to 7 cm), often grouped in bundles, can be observed even with the naked eye (V group: macroscopic fibres, after García-Romero and Suárez, 2013; Fig. 2).

(INSERT FIGURE 2)

These fibres are usually flexible (Fig. 2), with a cream to light brown colour, and readily split to thinner fibrils if manipulated. They usually crystallize parallel to the vein selvages (slip type), with thickness from < 1 mm to 1 cm. This morphology is common in the palygorskite-sepiolite group minerals, although massive, cardboard-like felts or ‘mountain leather’ (Grim, 1968; Frost *et al.*, 1998; Imai and Otsuka, 2000) of tangled fibres forming an apparently continuous mat are also typical. Observations under the stereo-microscope showed that these long macroscopic fibres are bundles composed of many thinner fibrils, intricately intergrown and separable by mechanic stress.

3.2 Scanning, Transmission and Analytical Electron microscopy

SEM investigation showed that the bundles observed at the macroscopic scale or by optical stereomicroscope are composed of entangled, elongated and mainly parallel thinner fibrils, < 0,1 μm width and > 150 μm long (IV group: very long fibres, García-

Romero and Suárez, 2013). When mechanically stressed, these thinner units show great flexibility (Fig. 3).

(INSERT FIGURE 3)

Owing to their sensitivity to the electron beam after a few seconds of exposure these fibres degrade by losing crystallinity and habit. Thus, high-magnification images were impossible to obtain by TEM. SAED images from fibre sections showed rings relating to the different relative rotations of the fibres along their axis lengths. The brightness of the diffraction spots comes out in favour of at least a medium crystallinity degree, although other experimental parameters may also play a role. Low magnification TEM images of the sample thinned perpendicular to [001] (lengthening axis of the fibre bundle) show a large open texture, with spaces among the fibre bundles and sometimes even among the single primary units – or laths (interfibre or open porosity: García-Romero and Suárez, 2013; Fig. 4.a).

(INSERT FIGURE 4)

Fibre sections cut perpendicular to the bundle axis show nearly regular rhomboidal (thinner fibres) to parallelogram-like cross-sections (larger units, ‘parallelograms’ hereafter). Similarly oriented parallelograms are grouped in clusters; separate clusters show different orientations (Fig. 4.b). The parallelogram dimensions vary approximately between 150-750 (base) and 80-250 Å (height). Parallelograms with dimensions larger than those given above show an incipient ‘opening’ process (fibre separation), similar to the defibrillation observed in several sections of amphibole asbestos (e.g. Gunter *et al.*, 2007). This ‘opening’ process causes fragmentation of larger into smaller parallelograms and occurs mainly along the traces of the (110) preferential cleavage plane (dashed white lines in Fig. 4.b). The separation starts mainly

from the fibre rim, deepening inside the core along a crankshaft direction which forms near 120° and 60° angles (Fig. 5). These parallelograms are not tightly aggregated as the global texture is characterized by medium to large interstitial spaces, sometimes larger than the parallelograms themselves, filled by an almost continuous thin film which, in some cases, apparently link the fibres. The film could not be characterized by electron probe microanalysis because of the very low analytical electron microscopy X-ray counts. This result suggests the presence of possible organic material, an assumption supported by FT-IR investigation (see paragraph 3.4). The film, located even in the smaller interstices (indicated by arrows in Fig. 4.b) suggests that it may be involved in the ‘opening’ process, favouring defibrillation and fibre separation of larger bundles into thinner fibrils (rods and/or laths: García-Romero and Suárez, 2013). These interstitial spaces, however, are not a consequence of beam damage to the sepiolite crystals because the parallelogram edges are clearly defined. A similar mechanism was recently described by Sciré et al. (2011), who observed how aliphatic hydrocarbons associated to flexible fibrous phyllosilicates (such as “serpentine”) trigger defibrillation of bundles into primary units, producing an atypical foam-like texture.

(INSERT FIGURE 5)

TEM observations performed on sections parallel to the fibre axis are less significant, owing to the large number of fibrils along the thickness of the section (consequent to their small width) and their rapid damage under the electron beam. This observation indicates also the high degree of parallelism among fibrils.

The open channel defects (OCD) described by Krekeler and Guggenheim (2009) in a sepiolite specimen from Helsinki are less common here. These defects involve the

omission of single to multiple polysomes in a fibre (commonly four); sometimes multiple OCD appear in the same fibre, especially in the larger units (Fig. 4.b).

The chemical composition measured by SEM-EDS (Table 1) is consistent with previously published data (Suàrez and Garcia-Romero, 2011) and shows Si and Mg as the main components (as well as O and H), together with small amounts of Fe and Al substituting for Mg in octahedral coordination; Ca was also sporadically detected.

(INSERT TABLE 1)

On an anhydrous basis, the following crystal chemical formula was obtained: $\text{Si}_{12,18}\text{O}_{32}(\text{Mg}_{7,18},\text{Fe}_{0,29})_{\Sigma 7,47}$ (total iron as Fe^{3+}). The slight Si excess in tetrahedrons with respect to the ideal value (12) may be a result of small disseminated quantities of amorphous silica, sometimes reported in literature (Karakaya et al., 2011). This silica, undetected by XRPD and FT-IR, is probably a result of excess SiO_2 content dissolved in the circulating low- T hydrothermal fluids which presumably precipitated this sepiolite (see paragraph 3.6). A low octahedral content (i.e. < 8) is not uncommon in sepiolite owing to any trivalent iron and vacancies (Suàrez and Garcia-Romero, 2011).

3.3 Thermogravimetric analysis

(INSERT FIGURE 6)

Thermogravimetric (TGA) and heat flow (DSC) data collected on the Perletoa sepiolite (Fig. 6) are consistent with previous works (Nagy and Bradley, 1955; Brauner and Preisinger, 1956; Martin Vivaldi and Cano Ruiz, 1956; Caillère and Hénin, 1957; Preisinger, 1959; Hayashi *et al.*, 1969; Nagata *et al.*, 1974; Rautureau and Mifsud, 1977; Ruiz *et al.*, 1996; Weir *et al.*, 2002; Hubbard *et al.*, 2003; Ovarlez *et al.*, 2006,

2009). Although in sepiolite correlations between TGA features and structural changes are not straightforward nor partitioning among different kinds of water so strict (Mifsud *et al.*, 1987), reasonable interpretations can be sketched. All main events and related attributions are detailed in Table 2.

(INSERT TABLE 2)

The weight loss recorded between room T and 110 °C (6.5 %), related to the release of superficially adsorbed water and less severely bound zeolitic H₂O, is slightly lower than that recorded in literature (i.e. about 10-11 %: Jones and Galan, 1988; Frost and Ding, 2003). This deficiency affects the hygroscopic rather than zeolitic H₂O content, as the roughly continuous organic film covering the fibres surface (see paragraph 3.2) probably limits the adsorption of physisorbed water. Loss of the 2nd half of structural OH₂, extending from 320 to 620 °C, is marked by no apparent signal in the derivative weight nor heat flow profiles (contrarily to Martin Vivaldi and Fenoll Hach-Ali, 1970 and Giustetto *et al.*, 2011c) thus certifying the slow progression of such an event.

Combustion of hydrocarbons should give rise to an exothermic peak around 500° C (Sciré *et al.*, 2011). Although a careful examination of the DSC profile reveals a weak exothermic hump around 470-480° C, no such an event is clearly observed here possibly due to the limited amounts of organic material forming the coating. The overall recorded weight loss (\cong 16.5 %) is slightly lower than that of other sepiolites (i.e. 18.5-19.5 %: Frost *et al.*, 2009; Giustetto *et al.*, 2011c), possibly as the result of the ascertained deficiencies in the hygroscopic H₂O content.

3.4 Fourier Transform IR spectroscopy

(INSERT FIGURE 7)

FT-IR spectra were collected on the Perletoa sepiolite in air at room T and in vacuum ($P < 5 \cdot 10^{-4}$ mbar) upon heating at 120 and 150 °C, to eliminate contribution of the broad signals related to zeolitic H₂O. Data are consistent with previous works (Cannings, 1968; Hayashi *et al.*, 1969; Mendelovici, 1973; Nagata *et al.*, 1974; Prost, 1975; Mendelovici and Portillo, 1976; Post, 1978; Myriam *et al.*, 1998; Frost *et al.*, 2001; Jung and Grange, 2004; Ovarlez *et al.*, 2009; Giustetto *et al.*, 2010; Giustetto *et al.*, 2011b; Bukas *et al.*, 2013). Attribution of vibrational modes and related evolution with T rise are summarized in Table 3.

(INSERT TABLE 3)

The FT-IR spectrum collected in air at room T (pattern 1 in Figg. 7 and 8) is dominated by the broad signals related to both physisorbed and zeolitic H₂O, which mask the narrower bands of hydroxyls having stronger interactions. Weak but clearly visible vibrational modes can be appreciated in the 1350-1550 cm⁻¹ region (at 1384 and 1509 cm⁻¹, separated by a broader hump at 1444 cm⁻¹; pattern 1 in Fig. 8, magnification; Table 3), where no bands of sepiolite should appear. These bands are consistent with those sometimes observed in literature on gabbroid or serpentinized/carbonated mantle-derived ultramafic xenoliths as well as saponite-rich clays originated from hydrothermal serpentinitic systems (Ciliberto *et al.*, 2009; Sciré *et al.*, 2011; Manuella *et al.*, 2012) and attributed to $\delta(\text{C-H})$ of saturated aliphatic hydrocarbons. Their presence can therefore be related to the organic film which fills the interstices among different fibres, possibly inducing the ‘opening’ of bigger parallelograms into thinner units as hinted by TEM (see Fig. 4.b, paragraph 3.2).

(INSERT FIGURE 8)

Evacuation at room T (pattern 2 in Figg. 7 and 8) causes loss of physisorbed and most zeolitic H_2O with disappearance of all related modes; conversely, all structural OH_2 is still preserved. This allows detection of weaker features at 3205, 2960, 2930 and 2854 cm^{-1} which represent the counterpart, in the stretching region, of those C–H modes related to aliphatic hydrocarbons (Silverstein *et al.*, 2005; Sciré *et al.*, 2011). The high intensity ratio between the 2930-2854 cm^{-1} signals [asymmetric and symmetric stretching of methylene (CH_2) groups respectively] and that at 2960 cm^{-1} [asymmetric stretching of methyl (CH_3) groups] indicates presence of hydrocarbons with long aliphatic chains (Coelho *et al.*, 2006).

When evacuation is coupled to temperature rise, release of OH_2 is due to happen at lower T than in air. In the adopted vacuum conditions, loss of the 1st OH_2 half starts at 120 °C and proceeds until 175-200 °C (Serna *et al.*, 1975; Giustetto *et al.*, 2011b), causing structure folding (Preisinger, 1963; Serna *et al.*, 1975, Post *et al.*, 2007), reduction of the tunnel width and transformation from tetra- to di-hydrated sepiolite ($\text{SEP}_4\text{H}_2\text{O}$ and $\text{SEP}_2\text{H}_2\text{O}$ respectively; Ovarlez *et al.*, 2011). Recent studies showed that these processes can be affected by incorporation of guest molecules in the tunnels, which prevents structure folding (Ovarlez *et al.*, 2009, 2011; Giustetto *et al.*, 2011b, 2012). Presence of the 3680 and 3674 cm^{-1} modes in the outgassed spectrum at 150 °C (pattern 4 in Fig. 7; Table 3) accounts for coexistence of folded and unfolded portions of the structure (Jung and Grange, 2004; Ovarlez *et al.*, 2009). Furthermore, a broad signal appears at 3423 cm^{-1} which merits further discussion. A similar mode (3420 cm^{-1}) was reported by Giustetto *et al.* (2011b) in a sepiolite + indigo (2 wt %) composite heated at 150 °C in vacuum, and attributed to structural OH_2 perturbed by H-bonds with the incorporated dye. In the present study, the band tentatively result from unspecific

interactions existing between structural OH₂ at the tunnel edges and superficial grooves (Benli *et al.*, 2012) and the almost continuous hydrocarbons film covering most fibrils. This suggestion is supported by the a similar signal in spectra collected on composite films between xylan-type hemicelluloses (polysaccharides) and fibrous sepiolite, indicative of H-bonding interactions (Sárosy *et al.*, 2012).

Bands related to the hydrocarbons film covering the sepiolite fibres are basically unaltered by *T* rise, but tend to sharpen (curves 3 and 4 in Fig. 7). In the bending region a broad hump is observed at 1455 cm⁻¹ bounded by sharper signals at 1509 and 1384 cm⁻¹ (Fig. 8, magnification; Table 3).

3.5. Crystal structure refinement

Despite all inevitable limits due to fine grain size, modest crystallinity and fibrous habit, which cause the refinement of this highly defective clay mineral to be troublesome (Guggenheim and Krekeler, 2011), a structural model for the Perletoa sepiolite was proposed using the Rietveld method on powder synchrotron XRD data consistently with previous studies (Post *et al.*, 2007; Giustetto *et al.*, 2011a). Recently pioneering single-crystal XRD data were presented by Sanchez del Rio *et al.* (2011).

Initial fractional coordinates were taken from Giustetto *et al.* (2011a); only Mg was located in octahedral sites and small quantities of Ca in the tunnels were neglected. The low-angular position and asymmetry of the strong (110) reflection ($d_{110} \cong 12.2 \text{ \AA}$), bringing information about the channel content (Ovarlez *et al.*, 2009), prevent its adequate fit so that in Rietveld procedures it is generally excluded (Chiari *et al.*, 2003; Post *et al.*, 2007; Post and Heaney, 2008; Giustetto *et al.*, 2011a). The refinement was

first carried out in the $1.5\text{--}41.5^\circ 2\theta$ range, including the strong (110) reflection, which was later excluded by maintaining the structural model refined so far.

(INSERT FIGURE 9)

(INSERT TABLE 4)

Fig. 9 shows the observed and calculated XRPD patterns of the Perletoa sepiolite and the related difference curve [magnification: patterns after exclusion of the (110) reflection]. Final refinement data, agreement factors and unit-cell parameters are listed in Table 4. Fractional coordinates, occupancy factors and displacement parameters for all atoms are reported in Table 5; the refined model is shown in Fig. 10.

(INSERT TABLE 5)

(INSERT FIGURE 10)

The goodness of fit is inevitably influenced by the disorder and multiple defects typical of these clay minerals (Giustetto and Chiari, 2004; Krekeler and Guggenheim, 2009), but the quality of the refined model is consistent with previous works (Artioli and Galli, 1994; Chiari *et al.*, 2003; Post *et al.*, 2007; Post and Heaney, 2008, Giustetto *et al.*, 2011a). Although the relatively high standard deviations prevent an accurate discussion of the finer structural details, the unit cell shows an increase ($0.03\text{--}0.05 \text{ \AA}$) in the length of all parameters with respect to previous models (i.e. Post *et al.*, 2007; Giustetto *et al.*, 2011a), mostly along y , with consequent increase in the cell volume ($\cong 1 \%$). The number of zeolitic H_2O molecules per unit cell (15.97) is in agreement with the ideal formula (16; Preisinger, 1959). Despite uncertainties remain about their sharp locations, these molecules are in sites consistent with those of Giustetto *et al.* (2011a); mutual $\text{O}\cdots\text{O}$ distances imply existence of a H-bond network (Jeffrey, 1997). The low

occupancy of ZW14 (Table 5) suggests that this molecule may alternatively occupy two close and equivalent sites.

3.6. *Origin of aliphatic hydrocarbons and sepiolite genesis*

The presence of hydrocarbons hosted in mafic to ultramafic, serpentinitic or gabbroic complexes involved in hydrothermal systems is well-known in literature (Charlou *et al.*, 1998; Sciré *et al.*, 2011). Hydrocarbons genesis can be either biotic or abiotic: the former involves biological processes including bacteriogenesis and thermogenesis (Schoell, 1988) whereas the latter is attributed to the Fischer-Tropsch-type (FT-t) reaction occurring in serpentinite-hosted hydrothermal systems (MacDonald and Fyfe, 1985; Konn *et al.*, 2009). The FT-t reaction is an exothermic reduction of CO and CO₂ by gaseous H₂ catalyzed by group VIII metal ions (such as Fe, Co, Ni) or their oxides. The reactant C oxides mainly derive from mantle CO₂ dissolved in hydrothermal fluids (Charlou *et al.*, 2002), whose concentration range from 3 to 20 mM. Furthermore, according to Ciliberto *et al.* (2009) CO₂ may be liberated during serpentinization of peridotitic olivine. In contrast, gaseous H₂ may be derived from the hydration of primary mafic minerals of peridotites (Marcaillou *et al.*, 2011). Initially the FT-t reaction implies dissociation of the adsorbed CO with formation of an intermediate carbide, whose presence was detected in the early serpentinization products of peridotites (Pikovskii *et al.*, 2004). Such an intermediate compound reacts with dissociated H₂ thus forming methane or higher hydrocarbons via insertion of –CH₂– monomers in a growing Fischer-Tropsch chain (Schultz, 1999).

The collection site of the studied sepiolite specimen plots in the area of the Combin Zone, formed by dominant calcshists which locally contain clasts of metabasites, Mn-rich quartzites and massive to foliated serpentinite bodies locally occurring as metabreccias containing magnetite- or carbonate-rich levels (Gasco and Gattiglio, 2011). The Combin Zone is one of the two units – along with the Zermatt-Saas Zone – belonging to the Piedmont Zone which is part of the Piedmont-Ligurian basin of the Neo-Tethys ocean (Mahlen *et al.*, 2005). This ocean spread during the Late Jurassic-Cretaceous (Piccardo *et al.*, 2001), when its closure started and completed during the early Tertiary as a result of the European and African plates collision.

The Piedmont-Ligurian oceanic basin is represented by ophiolites exposed along the Western Alpine-Northern Appennine orogenic chain (Piccardo *et al.*, 2001). A series of petrologic evidences supports the strong similarity of the Neo-Tethys ocean with ultra-spreading oceanic basins (e.g., Bach and Früh-Green, 2010) rather than with the traditional Penrose ophiolite model (Lagabrielle and Lemoine, 1997), such as:

- i) the abundance of serpentinitized spinel lherzolites intruded by gabbros;
- ii) the MORB affinity of gabbros and basaltic vulcanites (Piccardo *et al.*, 2001; Piccardo, 2008);
- iii) the lack of sheeted dykes and gabbroic layer 3 and the very low volumes of gabbros and basalts.

Furthermore, Tethys ophiolites bear mineralogical assemblages attributed to hydrothermal processes (Manatschal and Müntener, 2009), including high-temperature (HT) hydrothermal alteration observed in mylonitized gabbros and serpentinitization with formation of opicalcites in serpentinitized peridotites. Both processes resemble those observed in modern slow-spreading oceanic basin (e.g., Mid-Atlantic Ridge; Boschi *et*

al., 2006). Serpentinization primarily involves the complete alteration of olivine grains, whereas ortho- and clinopyroxenes are only occasionally preserved (Manatschal and Müntener, 2009). Ophicalcites, representing the *in situ* replacement of serpentine by calcite (Klein and Garrido, 2011), consist of clast- or matrix-supported breccias (Manatschal and Müntener, 2009) in which fragments of serpentinized peridotites are immersed in a matrix of red limestones and/or cemented by sparry calcite. The presence of ophicalcites provides evidence for hydrothermal activity at the Neo-Tethys seafloor at temperatures of 100-150 °C (Früh-Green *et al.*, 1990). Carbonate precipitation occurs in modern seafloor due to circulation of alkaline (pH = 9-11) and Ca-rich fluids in serpentinite-hosted hydrothermal systems (Schroeder *et al.*, 2002; Ribeiro da Costa *et al.*, 2008; Bach and Früh-Green, 2010; Lavoie and Chi, 2010) or by oxidation of abiotic methane (Schwarzenbach *et al.*, 2012). In addition, the finding of Mn-rich quartzites in the Tethys ophiolites has been interpreted by Tumiati *et al.* (2010) as a clear evidence for an oceanic hydrothermal origin of manganese ore deposits.

Serpentinite-hosted hydrothermal systems – analogous to those of the Combin Zone where the investigated sepiolite was collected – are ideal sites for the production of abiogenic hydrocarbons via the FT-t reaction, by means of progressive polymerization and polycondensation reactions (Konn *et al.*, 2009; Taran *et al.*, 2007). Since a widely accepted hypothesis for sepiolite genesis involves direct precipitation from an aqueous solution saturated with Mg and silica (Jones and Galán, 1988; Galán and Pozo, 2011) and with an insignificant Al activity (Birsoy, 2002), the Perletoa sepiolite probably originated from a similar process.

The absence of evident dolostones or dolomitic limestones sources in the area under study suggests that aqueous, low temperature (150-300 °C) hydrothermal fluids

probably derived Mg required to produce sepiolite by the weathering of surrounding serpentized ultramafic rocks belonging to the nearby ocean-derived Piemonte Zone (Fig. 1). Si, Fe and Al were possibly provided by circulation of fluids extracted from serpentinites, as due to serpentization of olivine and clinopyroxene (Augustin *et al.*, 2008). Very small quantities of amorphous silica, whose presence can be assumed following crystal chemical formula calculations (see paragraph 3.2) and possibly due to a SiO₂ excess in these aqueous solutions, are expected to favour sepiolite precipitation (Birsoy, 2002).

The intimate association of the Perletoa sepiolite fibres with their aliphatic hydrocarbons sheaths of presumed abiogenic origin (an occurrence never reported before in literature) has therefore to be considered as the result of a complex hydrothermal process, which possibly generated both components in separate but sequential steps. The sheath of hydrocarbons, which occupies even the smaller interstitial spaces, limits the amount of adsorbable hygroscopic water and probably interacts with the fibres surface *via* specific bonds. These interactions, both mechanical and chemical, are expected to play a key-role in favouring defibrillation ('opening' process schematized in Fig. 5) of thicker bundles into thinner fibrous units (rods and/or laths). Consistently with the chronology proposed by Sciré *et al.* (2011) formation of sepiolite probably came first, due to crystal growth through the oriented aggregation of smaller subunits (laths) precipitated from Si/Mg rich fluids which gradually grouped into thicker bundles (García-Romero and Suárez, 2013). The abiotic genesis of hydrocarbons *via* FT-t reaction started later, thus allowing the covering of the bundles with a thin organic sheath. Mutual interactions between the hydrocarbon coating and the sepiolite fibres surface slowly but inexorably opened new interstices among the

different units, exposing new surfaces and promoting the beginning of the defibrillation process. The continuous production of hydrocarbons brought new organic matter to further slip into the formerly opened spaces, thus endorsing reiteration of the mechanism until discharge of primary sepiolite laths was reached.

The described phenomenon shows several analogies with Sherman (1970), who patented a protocol for defibrillating asbestos. An insoluble ethylenically unsaturated monomer was added to primary activated fibres dispersed in water (pH 4-5); its consequent polymerization formed an *in situ* organic coating on the surface of the asbestos fibrils. Such a method was aimed at obtaining a large number of primary fibrils, each coated by a polymer film, thus facilitating their dispersion in synthetic plastic compositions and reducing the tendency of such materials to crack or degrade after asbestos addition and mixing. It is curious to notice how such an artificial process, devised more than 40 years ago, has been mimicked by nature on an analogous material (this sepiolite occurrence, a potential asbestos substitute: see e.g., Kavas *et al.*, 2004) by exploiting the coexistence of peculiar environmental conditions in a specific geological context.

4. Conclusions

The progressive defibrillation of the Perletoa sepiolite, triggered by the hydrocarbons sheath, not only causes a significant increase in the interfibre porosity (open texture) but also affects the fibre morphology enhancing its aspect ratio (length vs. thickness) from 'high' to 'very high'. The thinner and exceptionally long fibrils

(rods and/or laths), therefore, potentially become more dangerous for human health due to their carcinogenic potential if dispersed in air and breathed in high doses.

However, it cannot be excluded that the hydrocarbons sheath could affect the surface reactivity of the sepiolite fibrils altering their chances of interacting with the pulmonary epithelium. For example, only the breathing of quartz particles with fresh surfaces increases the risk of lung damages due to their enhanced reactivity, whereas old and contaminated grains appear to be less dangerous (thus explaining why miners tend to be affected most; Schins *et al.*, 2002; Albrecht *et al.*, 2005). Besides, the toxicity of occasional exposure to aliphatic hydrocarbons *per se* (inhaled, skin absorbed or ingested) has already been acknowledged in literature (Farinha *et al.*, 2011). Though the fibrous habit supposedly increases the risk, if the analogy holds then presence of this organic coating may enhance or else reduce this sepiolite noxiousness.

In the most recent report on the carcinogenic risks for humans caused by some silicates, the International Agency for Research on Cancer (IARC, 1997) stated that there is an “inadequate evidence in humans for the carcinogenicity of sepiolite”; despite this, a “limited evidence in experimental animals for the carcinogenicity of long sepiolite fibres” (length > 5 µm) was suggested. Fibres shorter than 5 µm, on the contrary, were considered harmless. On the whole, sepiolite fibres were included by IARC in “Group 3”, meaning that further investigations are needed to evaluate their possible carcinogenicity or noxiousness (IARC, 2012). However, no data on sepiolite carcinogenicity for humans were available to the IARC working group in 1997, when the report was published. Data on the toxic effects for humans were concerned mostly for fibres < 4 µm. Realistically, the possible health issues of fibres characterized by exceptional lengths, such as those observed here, are not known.

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Table Captions

Table 1. Chemical composition (weight % oxides and mean value) and cations number calculated on anhydrous basis (32 oxygens) of the studied Perletoa sepiolite.

Table 2. TGA weight losses and related attributions for the Perletoa sepiolite.

Table 3. FT-IR active vibrational modes and related attributions at different temperatures and vacuum conditions for the Perletoa sepiolite and the superficial hydrocarbons sheath.

Table 4. Crystal-chemical formula, space group, cell parameters and refinement data for the Perletoa sepiolite.

Table 5. Refined fractional atomic coordinates, occupancy factors and isotropic displacement parameters for the Perletoa sepiolite.

Tables

Oxides (wt%)	1	2	3	4	5	6	7	8	9	10	11	12	13	Average
SiO ₂	56.05	57.35	56.73	56.04	56.73	58.28	56.23	56.93	56.69	57.44	54.38	54.46	56.12	56(1)
Fe ₂ O ₃	1.41	1.32	1.97	2.01	1.94	2.33	2	1.62	1.98	2.11	1.52	1.44	1.71	1.8(3)
MgO	22.13	22.67	22.25	22.11	22.84	23.38	22.14	22.36	21.87	23.37	21.39	21.9	21.74	22.3(6)
CaO	0	0	0	0.74	0	0	0	0	0	0	0	0	0	=
Σ	79.58	81.34	80.95	80.91	81.51	83.99	80.38	80.9	80.53	82.91	77.29	77.8	79.57	81(2)
Cations														
Si	12.23	12.24	12.20	12.11	12.13	12.11	12.18	12.22	12.24	12.08	12.22	12.17	12.25	12.18(6)
Fe ³⁺	0.23	0.21	0.32	0.33	0.31	0.36	0.33	0.26	0.32	0.33	0.26	0.24	0.28	0.29(5)
Mg	7.20	7.21	7.13	7.12	7.28	7.24	7.15	7.16	7.04	7.33	7.17	7.30	7.07	7.19(9)
Σ	19.66	19.66	19.65	19.56	19.72	19.71	19.66	19.64	19.60	19.74	19.65	19.71	19.60	19.66(5)
Ca	0	0	0	0.17	0	0	0	0	0	0	0	0	0	=

Table 1.

Temperature range (° C)	Weight loss (%)	DSC event	Attribution and comments
25 – 110	6.5	endothermic (110° C)	Loss of physisorbed water and less severely bound zeolitic H ₂ O (tetra-hydrated sepiolite; SEP4H ₂ O)
120 – 310	3.5	endothermic (310° C)	Loss of residual zeolitic H ₂ O and 1 st fraction of structural OH ₂ (di-hydrated sepiolite; SEP2H ₂ O)
320 – 620	3.5	=	Loss of 2 nd fraction of structural OH ₂ (anhydrous sepiolite)
650 – 820	3.0	endothermic (810° C) endothermic (830° C) exothermic (850° C)	(Possible) decarbonation of calcite traces Loss of framework OH (dehydroxylation) and transformation to clinoenstatite (or an amorphous phase)
Total	16.5		

Table 2.

Sepiolite				
Room <i>T</i> in air (cm ⁻¹)	Room <i>T</i> in vacuum (5*10 ⁻⁴ mbar)	120° C in vacuum (5*10 ⁻⁴ mbar)	150° C in vacuum (5*10 ⁻⁴ mbar)	Attribution and comments
Stretching region (2750 - 3750 cm⁻¹)				
3246	=	=	=	v(OH) of physisorbed water, lost by evacuating at room <i>T</i>
3358	=	=	=	v(OH) of zeolitic H ₂ O, lost by evacuating at room <i>T</i>
=	=	=	3423	Unspecific interactions of structural OH ₂ (possibly with the aliphatic hydrocarbons sheath)
3568	3550	3550	3531	v(OH) of structural OH ₂ ; red-shift at 3550 and 3531 cm ⁻¹ are due to H-bond and symmetric v(OH ₂) variations
3623	3623	3623	3644-3603	v(OH) of structural OH ₂ ; splitting at 150° is due to variations in the antisymmetric and symmetric v(OH ₂)
3690	3680	3680	3680	v(OH) of hydroxyl in O sheet; typical of unfolded sepiolite structure
=	=	=	3674	v(OH) of hydroxyl in O sheet; accounts for intervened folding of the sepiolite structure
=	=	=	3692	H-H repulsion of framework hydroxyls (Mg-OH) due to approaching of residual OH ₂ dislocated by structural folding
=	3719	3719	3719	v(Si-OH); gradual intensity decay with <i>T</i> rise
=	=	3738	3738	v(OH); perturbation of superficial silanols; Si-OH liberation at the edge of the O ribbons due to structure folding
Bending region (1400 - 1800 cm⁻¹)				
1660	1623-1613	1623-1613	1625	δ(H ₂ O) phys./zeolitic (room <i>T</i>) - δ(OH ₂) (vacuum/heat); intensity decay with <i>T</i> rise accounts for gradual OH ₂ loss
1212	1193	1195	1198	δ(Si-O)
Aliphatic hydrocarbons				
Room <i>T</i> in air (cm ⁻¹)	Room <i>T</i> in vacuum (5*10 ⁻⁴ mbar)	120° C in vacuum (5*10 ⁻⁴ mbar)	150° C in vacuum (5*10 ⁻⁴ mbar)	Attribution and comments
Stretching region (2750 - 3750 cm⁻¹)				
=	3205	3205	3213	v(C-H); slightly blue-shifts at 150° C
=	2960	2960	2960	v(CH ₃)
=	2930	2930	2930	v(CH ₂)
=	2854	2854	2854	v(CH ₂)
Bending region (1400 - 1800 cm⁻¹)				
1384	1384	1384	1384	δ(CH ₃) symmetric
=	=	1403	1403	δ(C-H); appears at 120° C
1444	1444	1455	1455	δ(CH ₂) scissoring, δ(CH ₃) asymmetric; blue-shifts at 120° C
1509	1509	1509	1509	δ(C-H)

Table 3.

Formula	Si _{12,18} O ₃₂ (Mg _{7,18} ,Fe _{0,29}) _{Σ7,47}
Space group	<i>Pncn</i>
<i>a</i> (Å)	13.442(2)
<i>b</i> “	27.072(3)
<i>c</i> “	5.2926(6)
V (Å ³)	1925.9(4)
Δρ _{min} , Δρ _{max} (e/Å ³)	-0.300; 0.305
R	0.0202
R _{wp}	0.0271
R(F ²)	0.1260
reduced χ ²	8.420

Table 4.

Atom type	Label	<i>x</i>	<i>y</i>	<i>z</i>	Fraction	<i>U</i> _{iso}
Mg	Mg1	0	0.027(1)	0.25	1.00	0.008(2)
Mg	Mg2	0	0.089(1)	0.75	1.00	0.008(2)
Mg	Mg3	0	0.143(1)	0.25	1.00	0.008(2)
Mg	Mg4	0	0.204(1)	0.75	1.00	0.008(2)
O	O1	0.075(2)	0.024(1)	0.549(6)	1.00	0.016(3)
OH	O2	0.088(2)	0.082(2)	0.096(7)	1.00	0.016(3)
O	O3	0.076(2)	0.139(1)	0.588(6)	1.00	0.016(3)
Si	Si1	0.198(1)	0.0280(7)	0.578(4)	1.00	0.011(2)
Si	Si2	0.197(1)	0.1415(6)	0.582(4)	1.00	0.011(2)
Si	Si3	0.216(1)	0.1944(5)	0.080(4)	1.00	0.011(2)
O	O4	0.096(2)	0.204(1)	0.082(7)	1.00	0.016(3)
O	O5	0.245(2)	-0.001(1)	0.328(4)	1.00	0.016(3)
O	O6	0.238(2)	0.0849(8)	0.574(4)	1.00	0.016(3)
O	O7	0.246(2)	0.161(1)	0.317(5)	1.00	0.016(3)
O	O8	0.253(2)	0.171(1)	0.812(5)	1.00	0.016(3)
O	O9	0.25	0.25	0.137(8)	1.00	0.016(3)
OH ₂	SW10	0.078(2)	0.2553(8)	0.572(7)	1.03(6)	0.025(7)
H ₂ O	ZW11	0.574(4)	-0.091(3)	0.034(9)	0.94(7)	0.10(4)
H ₂ O	ZW12	0.5	0.186(4)	0.25	0.76(9)	0.10(4)
H ₂ O	ZW13	0.5	0.013(3)	0.25	0.98(9)	0.10(4)
H ₂ O	ZW14	0.50(1)	0.168(4)	0.74(2)	0.19(4)	0.10(4)

Table 5.

Figure captions

- Figure 1. Simplified tectonic sketch-map of the Western Alps. Helvetic Domain: Mont Blanc-Aiguilles-Rouges (MB); Penninic Domain: Grand Saint Bernard Zone (SB) and Internal Crystalline Massifs of Monte Rosa (MR), Gran Paradiso (GP), Dora-Maira (DM) and Valosio (V); Piemonte Zone of calcschists (light green) with meta-ophiolites (dark green): Lanzo Ultramafic Massif (LM); Austro-alpine Domain: Dent-Blanche nappe (DB), M. Emilius nappe (ME) and Sesia Zone (SZ); Southalpine Domain (SA); Nappe of Embrunais-Ubaye Flysch (EU); Canavese Line (CL); Sestri-Voltaggio Line (SVL); PF: Penninic Thrust Front. BAL = chrysotile asbestos Balangero mine (Province of Torino). The star indicates the recovery position of the studied sepiolite specimen, near Perletoa village.
- Figure 2. Visual appearance of the Perletoa sepiolite: (a) thick bundles of fibres still attached to the serpentine substrate and (b) isolated. Notice how the fibrous habit is well evident even at the macroscopic scale.
- Figure 3. Secondary electron SEM image showing several bundles of thinner fibrils orientated approximately in the same direction. The flexibility of the Perletoa sepiolite is evidenced by the accentuated bending shown by some fibres.
- Figure 4. TEM micrographs of sepiolite fibres in cross section, observed along the [001] direction. At medium magnification the fibrils show a rhomboidal to parallelogram-like contour and quite a large open texture (a). At high magnification the incipient ‘opening’ process undergone by the central thick fibre, which causes subdivision in thinner units, can be better appreciated (split surfaces shown as dashed white lines). A roughly continuous and scarcely electron-dense film (possibly organic matter) surrounding the fibre contours and filling the interstices is indicated by arrows (b).
- Figure 5. Simplified scheme of the ‘opening’ process which splits thick and elongated rhombic prismatic fibres into thinner ones. Preferential cleavage on the (110) crystal plane causes progressive fragmentation of a thicker sepiolite fibre (a) and consequent partition in some (4) thinner fibrils, maintaining the same rhomboidal contour and elongated in the [001] direction (b). Evolution of the same process causes the thinner fibrils to progressively get far from each other (c). The actual process, as a matter of fact, forms less regular smaller units. (100) and (010) faces, corresponding to minor cleavage planes, are also indicated for sake of clarity.
- Figure 6: Observed TGA/derivative weight and heat flow/derivative heat flow curves for the Perletoa sepiolite. Vertical scale for the TGA curve is weight loss %.

Figure 7. FTIR spectra in the stretching region of the Perletoa sepiolite in air (1) and evacuated at room temperature (2), 120 °C (3) and 150 °C (4) respectively. Spectra are shifted on the Y axis for sake of clarity.

Figure 8. FTIR spectra in the bending region of the Perletoa sepiolite in air (1) and evacuated at room temperature (2), 120 °C (3) and 150 °C (4) respectively. Spectra are shifted on the Y axis for sake of clarity.

Figure 9. Observed (crosses) and calculated (solid line) patterns for the Rietveld refinement of the Perletoa sepiolite, together with the related difference (lower line).

Figure 10. Refined structure of the Perletoa sepiolite as resulting from the Rietveld procedure (crystal cell outlined in solid bars; H atoms arbitrarily added to both zeolitic H₂O and structural OH₂).

Figures

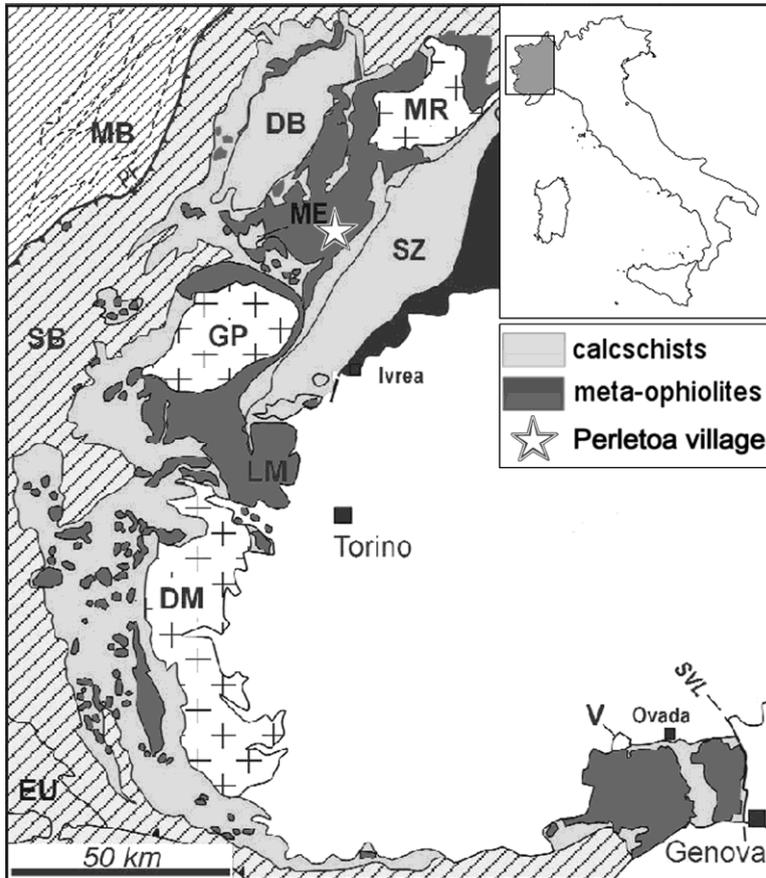


Fig. 1.

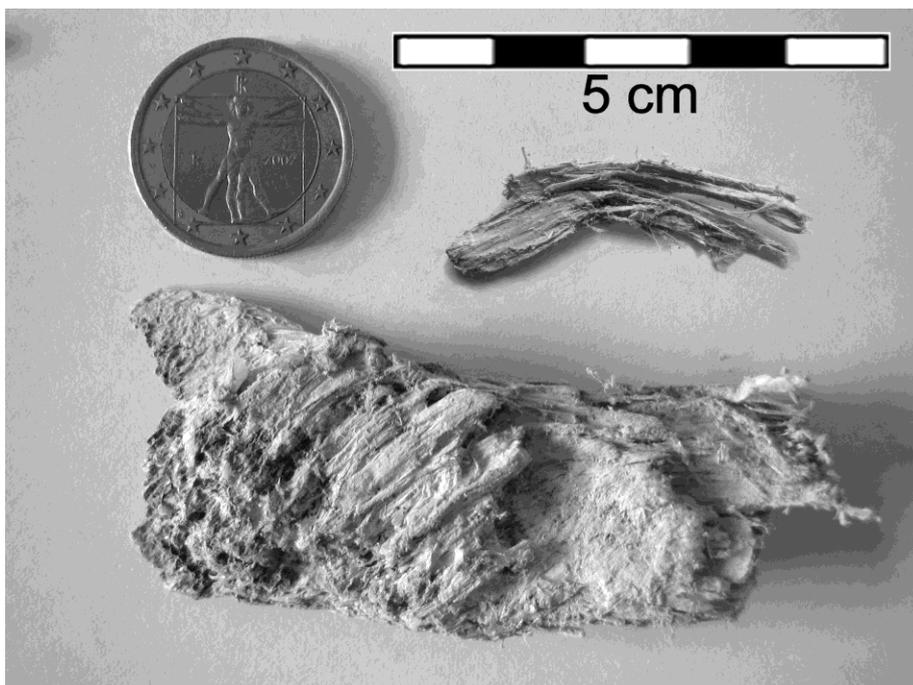


Fig. 2.

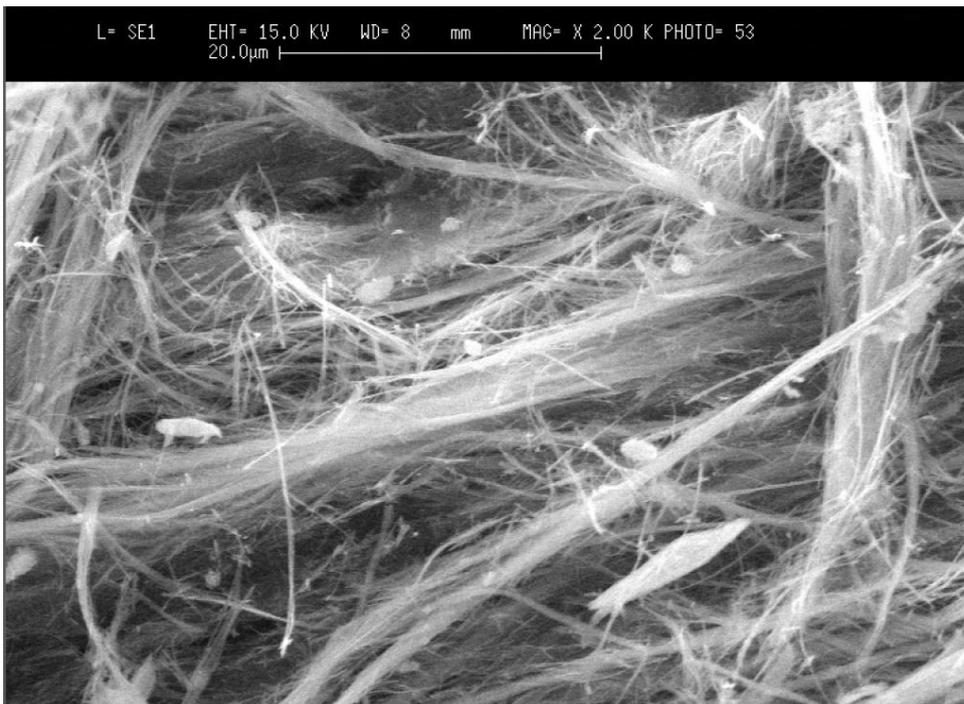


Fig 3.

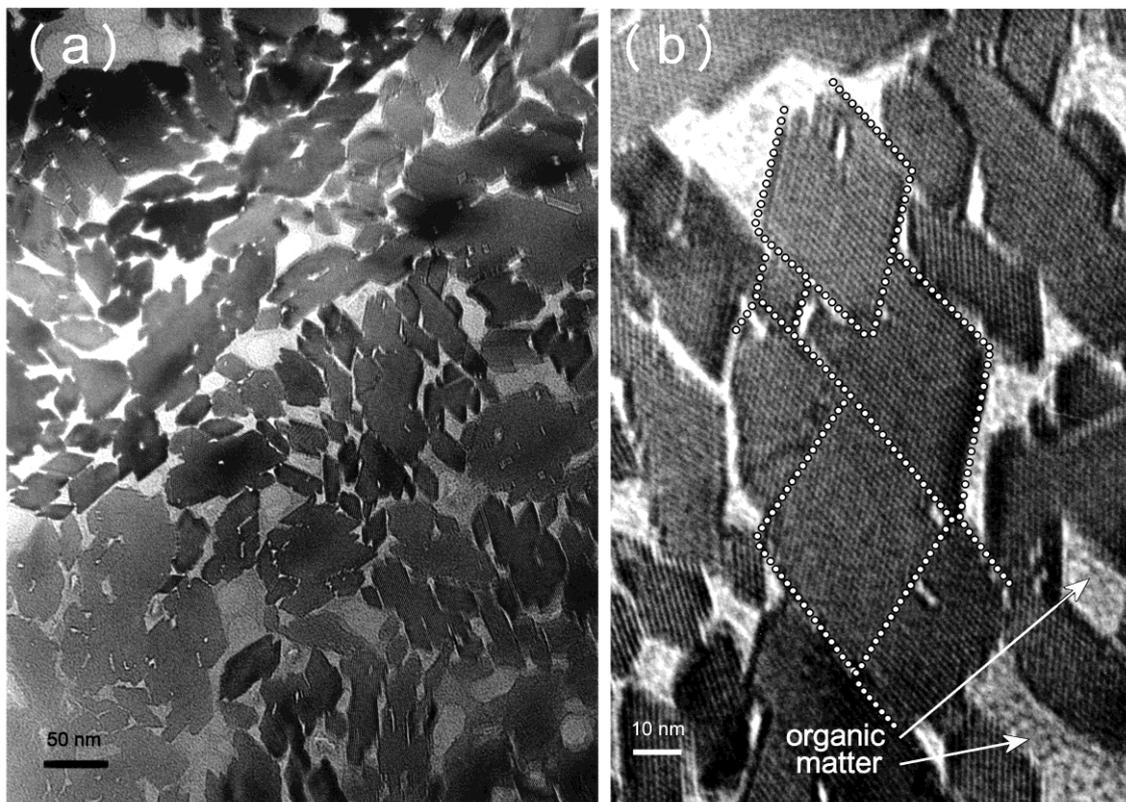


Fig. 4.

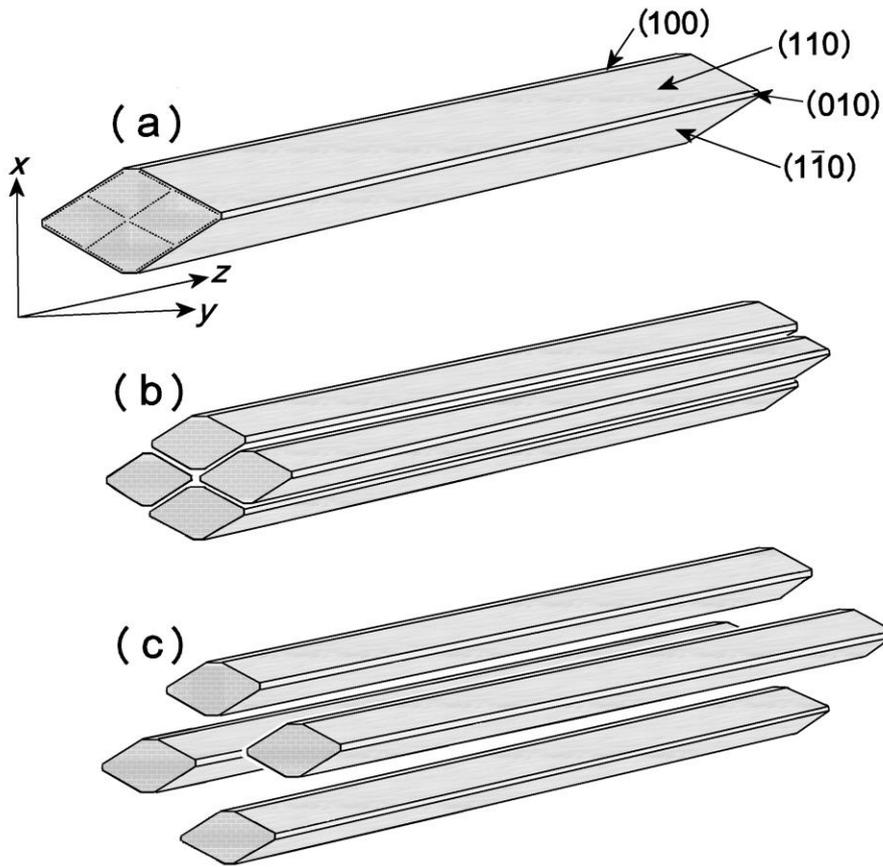


Fig. 5.

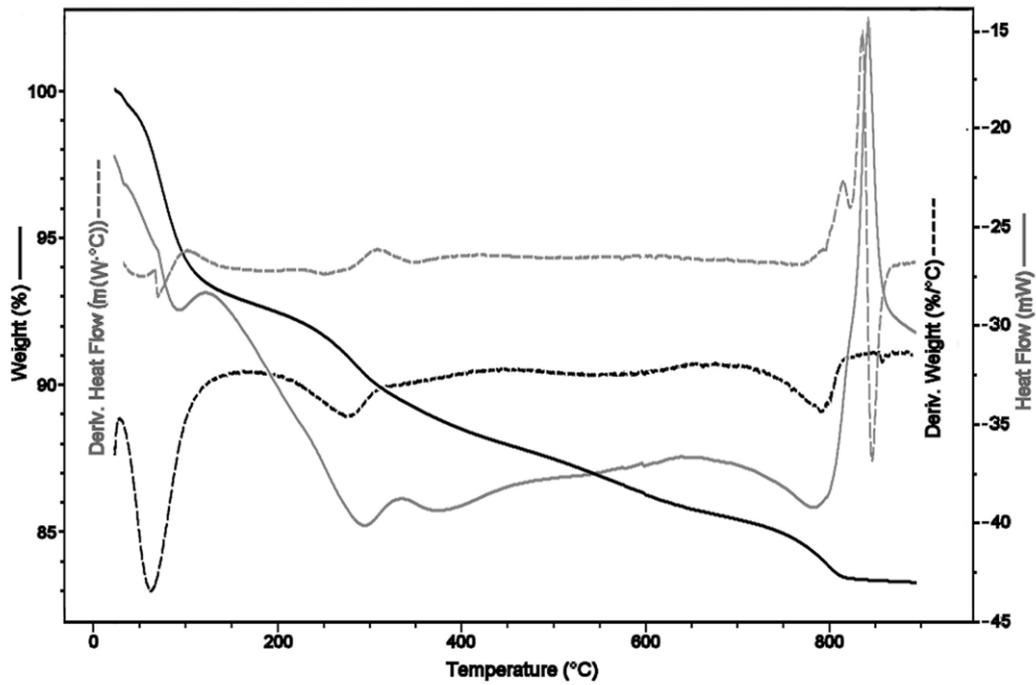


Fig. 6.

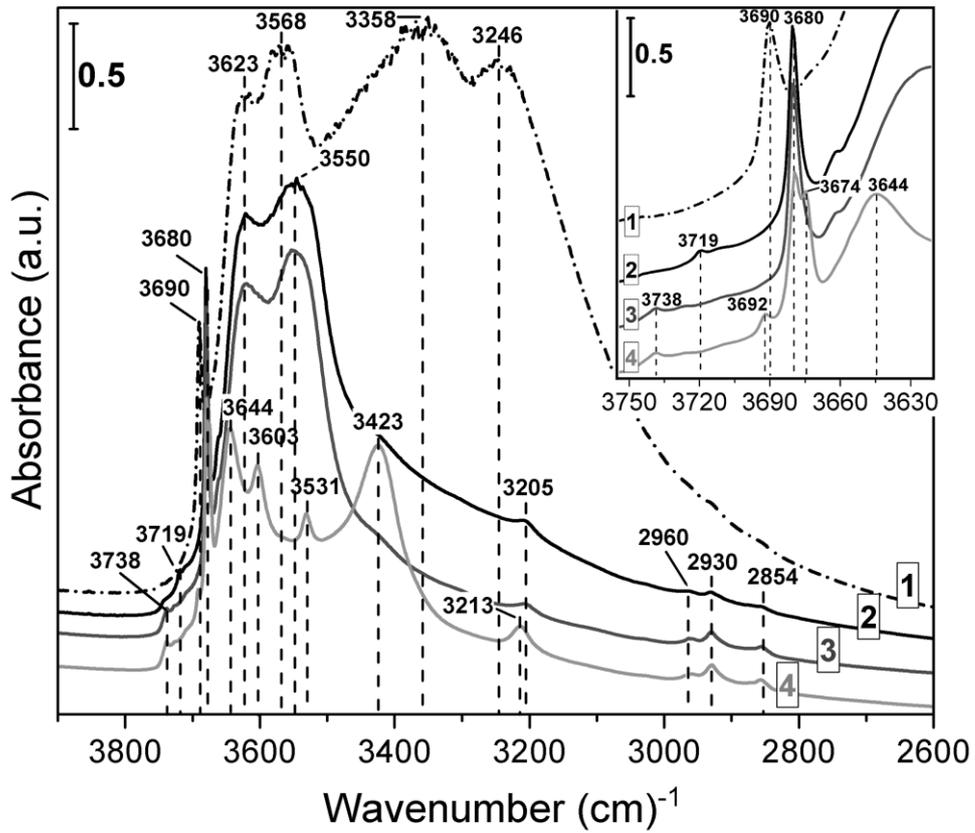


Fig. 7.

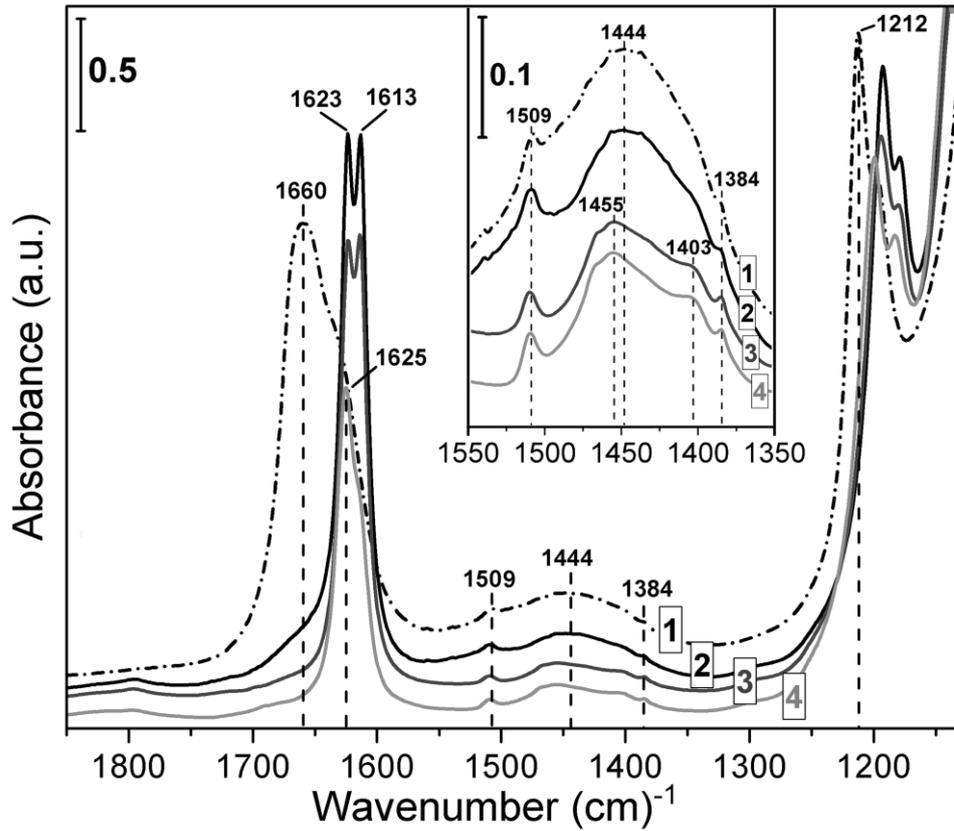


Fig. 8.

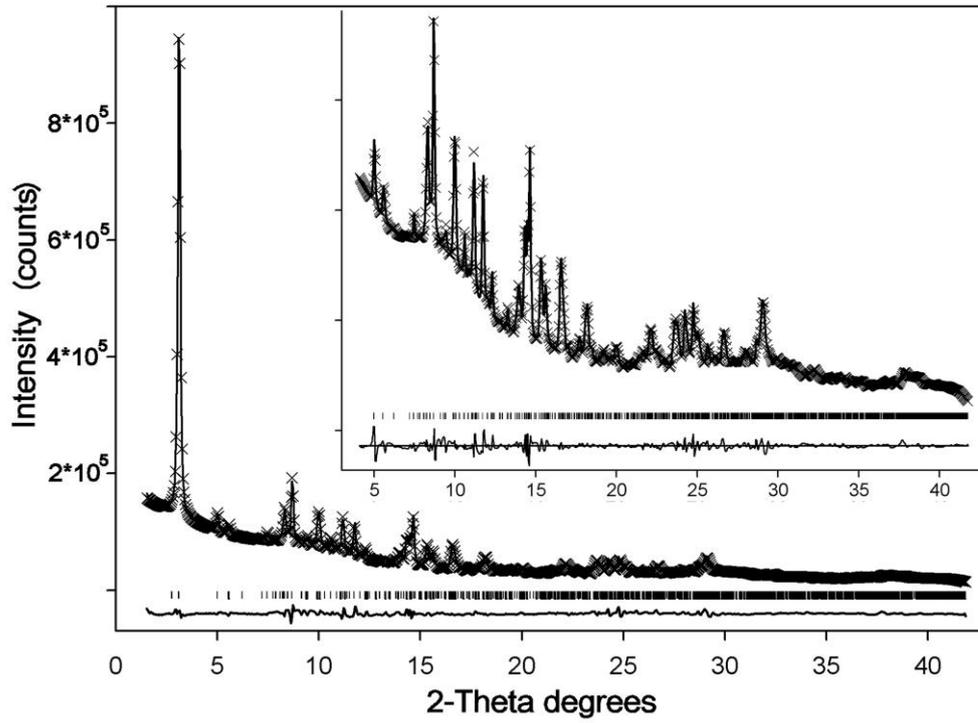


Fig. 9.

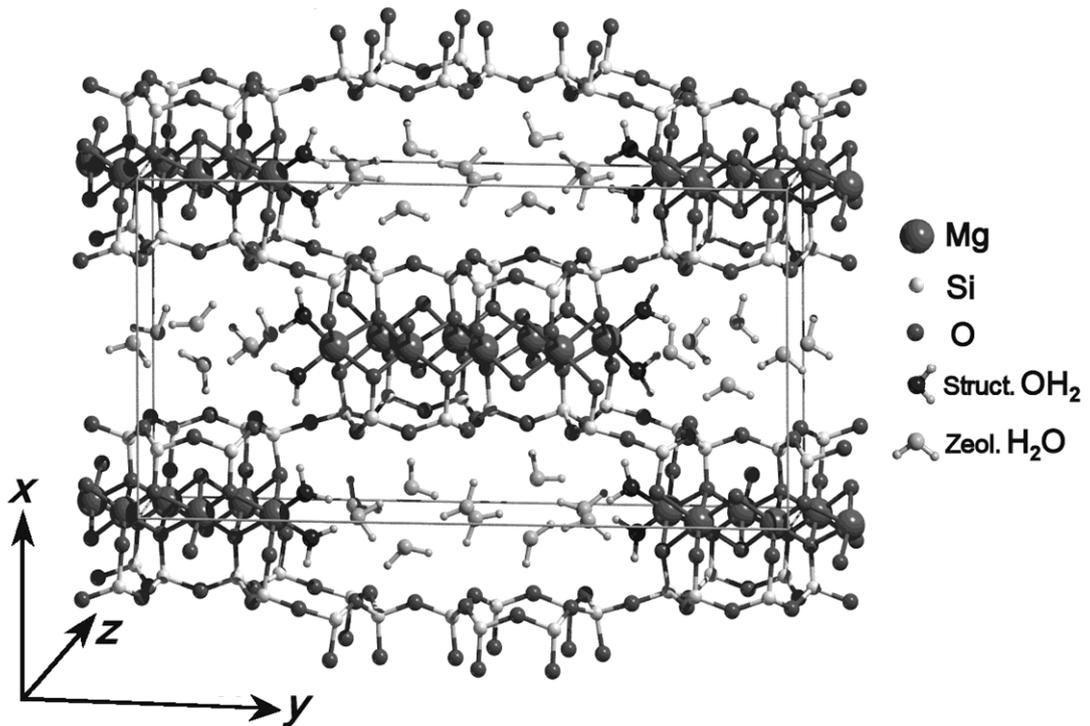


Fig. 10.