Biogenic calcium carbonate as evidence for life

Sara Ronca[1], Francesco Mura[2], Marco Brandano[1], Angela Cirigliano[3], Alessandro Grottoli[4], Massimo Reverberi[5], Danile Federico Maras[6], Rodolfo Negri[7], Ernesto Di Mauro[3] & Teresa Rinaldi[7]

5 [1]Department of Earth Sciences, Sapienza University of Rome, Italy
[2]CNIS-Center for Nanotechnology Applied to Industry of La Sapienza, 00185, Sapienza University of Rome, Rome, Italy
[3]Istituto di Biologia e Patologia Molecolari, CNR, Rome, Italy
[4]SARA EnviMob s.r.l., Rome, Italy
[5]Department of Environmental Biology, Sapienza University of Rome, Rome, Italy
10 [6]Soprintendenza Archeologia, Belle Arti e Paesaggio per la Provincia di Viterbo e l'Etruria Meridionale, Ministero della Cultura, Rome, Italy
[7] Department of Biology and Biotechnology, Sapienza University of Rome, Rome, Italy

Correspondence to: Teresa Rinaldi (teresa.rinaldi@uniroma1.it)

Abstract. The history of Earth is a story of co-evolution of minerals and microbes: not only numerous rocks arisen from life, but the life itself may have formed from rocks. To understand the strong association between microbes and inorganic substrates, we investigated the moonmilk, a speleothem of calcium carbonate of microbial origin, present in the Iron Age Etruscan Necropolis of Tarquinia, in Italy. These tombs present a unique environment where the hypogeal walls of the tombs are covered
20 by this speleothem. To study moonmilk formation, we investigated the bacterial community *in* the rock in which the tombs are carved: calcarenite and hybrid sandstone. We present the first evidence that moonmilk precipitation is driven by microbes *within* the rocks and not only *on* the rock surfaces. We also describe how the moonmilk produced within the rocks contributes to rock formation and evolution. The microbial communities of the calcarenite and hybrid sandstone displayed, at phylum level, the same microbial pattern of the moonmilk sampled from the walls of the hypogeal tombs, pointing out that the
25 moonmilk originates from the metabolism of endolytic bacterial community. The calcite speleothem moonmilk is the only known carbonate speleothem on Earth with undoubted biogenic origin, thus representing a robust and credible biosignature of life. Its presence in the inner parts of rocks adds to its characteristics as a biosignature.

1 Introduction

Whether other planets witnessed life like what is seen on Earth remains a complete mystery. The search for traces of extra-
30 terrestrial life suffers from the lack of durable and credible biosignatures. Some breakthroughs may happen soon from the current planetary explorations. The NASA Perseverance rover successfully landed on planet Mars in the Jezero Craters in 2021 and is collecting many specimens. Future missions are planned to retrieve those specimens, although not before 2031, hence opening the possibility to search for evidence of life from the first-ever samples returned from Mars. The European Space Agency (ESA) ExoMars programme is also planning to address the question of whether life has ever existed on Mars.

35   Earth has evolved through a long process of co-evolution between minerals and microbes (Cuadros, 2017; Cosmidis and
      Benzerara, 2022; Grosch et al., 2015; Hazen et al., 2008) and terrestrial rocks constitute an ideal system for the investigation
      of durable signs of life. Carbonates are good candidates as host-rock because, on Earth, they are principally of biogenic origin.
      Nevertheless, they did not receive the attention needed because of limited and sporadic evidence on Mars: lithologies revealed
      the presence of carbonates in the Nili Fossae Region (Ehlmann et al., 2008), at the Mars Phoenix landing site (Boynton, et al.,
40   2009), in the Columbia Hills of Gusev crater (Morris et al., 2010), and in deep rocks exposed by meteor impacts (Michalski et
      al., 2010). More recently, the analysis of weathering profiles revealed the widespread distribution on Mars of carbonates
      associated with hydrated minerals, providing evidence of past liquid water presence (Bultel et al., 2019). Carbonates were also
      detected in asteroids and meteorites, contributing to the understanding of the formation and evolution of our solar system
      (Pilorget et al., 2021; Kaplan et al., 2020; Voosen, 2020; Lee et al., 2014).

45   Carbonate rocks on Earth are of abiogenic or biogenic origin and occurred since the early Archean (> 3 giga annum, Ga) Eon
      when hydrothermal systems were ubiquitous. At that time, the carbonate rocks originated by massive carbonatization,
      silicification and potassium (K) (±Sodium) metasomatism of intermediate to ultramafic silicate precursors (Veizer et al., 1989).
      In contrast, Lower Archean marine carbonates are rare, they occur as very thin, discontinuous, and extensively mineralized
      beds generally replaced by chert as result of intense microbial iron (Fe) cycling (Pomar, 2020). The oldest known carbonate
50   production associated with biotic carbon dioxide (CO_2) sequestration are stromatolite structure 3.7 Ga old (Nutman et al.,
      2016). Successively, during Proterozoic (<2.4 Ga), shallow water carbonate production expanded favoring the development
      of carbonate platforms where abiotic and biotic carbonate precipitation took place (Grotzinger et al., 2000). The seawater was
      supersaturated with both calcite and aragonite as evidenced by the well preserved pseudomorphs of 'abiotic' aragonite and
      calcite (Pomar, 2020). According to Sumner and Grotzinger (1996), the rise in oxygen concentration at 2.2–1.9 Ga led to the
55   removal of Fe^{2+}, a strong calcite-precipitation inhibitor, from seawater and resulted in a shift from Archean to Proterozoic
      carbonates that are dominated by microbial activity (Pomar, 2020). Starting from the Cambrian Period, the benthic
      microbialites, the prime carbonate factory since the Late Archean, were still important but progressively decreased, while the
      biologically controlled carbonates appeared and expanded. The calcification of sessile, mostly colonial, metazoans and algae
      promoted the accumulation of biogenic carbonate sediments and the appearance and expansion of reefs (Pomar, 2020).

60   Carbonates are common constituents of the near-surface Earth crust, although carbonate phases may also occur deep in the
      mantle. They are compounds formed by the anionic complex, carbonate $(CO_3)^{2-}$ combined with metal ions such as calcium
      (Ca), magnesium (Mg), iron, manganese, sodium, barium, aluminium, zinc, copper, lead, uranium, or rare-earth elements.
      Uncommonly, carbonate phases are hydrated and contain hydroxyl or halogen ions or may include silicate, sulphate or
      phosphate radicals. Due to the large availability of Ca and Mg in crustal reservoirs (Hartmann et al., 2012), the $CaCO_3$
65   polymorphs, calcite and aragonite, and dolomite $(CaMg(CO_3)_2)$ are the most widespread carbonate minerals whose formation
      on Earth near-surface environments is widely related to biogenic or bio-mediated processes (Görgen et al., 2021).

      Carbonate biomineralization results in the formation of several mineral phases, the most common of which are: the $CaCO_3$
      anhydrous polymorphs calcite, aragonite and vaterite, the last being a metastable transitional phase, the hydrated forms

monohydrocalcite ($CaCO_3 \cdot H_2O$) and ikaite ($CaCO_3 \cdot 6H_2O$), and various amorphous phases (ACC). Moreover, in specific

70   environments (saline lakes, coastal lagoons) (Kaczmarek et al., 2017; Diloreto et al., 2021), microbial activity may promote

the formation of dolomite (ordered phase $CaMg(CO_3)$ by passing through the precursor phases of high Mg-calcite (disordered

4–36 mol % $MgCO_3$), disordered dolomite (disordered >36 mol % $MgCO_3$) and proto-dolomite (weakly ordered >36 mol %

$MgCO_3$). Carbonate mineral formation seems to proceed from amorphous or disordered phases towards the more stable and

ordered forms (Asta et al., 2020); crystal growth and morphologies are controlled by the medium composition, the microbial

75   extracellular polymeric substances (EPS), the Mg/Ca ratio and the presence of other ions. Thus, the mineral phases resulting

from abiogenic or biogenic activity are indistinguishable, and the identification of irrefutable biosignature, evidence of past or

present life, is still lacking (Changela et al., 2021; Javaux, 2019), hence, its necessity. Carbonate rocks, being mostly of

biogenic origin, could be considered in this perspective, but even if specific analyses could discriminate between biotic and

abiotic carbonate rocks (Blanco et al., 2013), the attempt to unequivocally distinguish between carbonates of biogenic or

80   abiogenic origin remains vacuous, especially in a search for evidence for life on other planets. So far, the presence of organic

materials in carbonate has not been an incontrovertible indicator of biogenicity (Berg et al., 2014).

Potential interest on the calcium carbonate minerals as possible biosignatures can be found by the study of a secondary calcite

deposit (speleothem), called moonmilk, formed by nanofibers of calcite (Borsato et al., 2000), commonly found on karst caves

surfaces. The moonmilk precipitation is promoted by the metabolic activities of a microbial community living in environments

85   with high calcium content (Banks et al., 2010; Cailleau et al., 2009; Cirigliano et al., 2018; Portillo et al., 2011). Nevertheless,

direct evidence that bacteria promote the precipitation of these nanofibers of calcite was still lacking. This evidence is essential

to define the moonmilk as a *bona fide* biosignature.

Recently, the moonmilk speleothem was discovered in the hypogeal ancient Etruscan tombs of the Monterozzi Necropolis

(Tarquinia, Central Italy) (Cirigliano et al., 2018; Tomassetti et al., 2017). This finding provides a unique opportunity to

90   compare the moonmilk which covers the walls and ceilings collected from 12 tombs excavated in two types of rock, the

calcarenite and hybrid sandstone. We provided insight into the formation of moonmilk, which can occur rapidly, i.e. between

10 and 50 years, and we reported that this speleothem originates from, and harbours a highly active microbial community

(Cirigliano et al., 2021a; Cirigliano et al, 2021b). Thus, the moonmilk formation requires the presence of microorganisms and

it is formed by nanofibers of calcite, a less common crystal habit for calcium carbonate, suggesting that this speleothem could

95   be proposed as a sign of life. In order to consider the moonmilk as a biosignature, one needs to find a strong association with

the rocks from which it is originated, i.e. the moonmilk deposition should be a geological process, and not just a phenomenon

of biomineralization in which the microorganisms use the rocks as a physical substrate. Hence, the need of a deeper analysis

of the mineralogy and of the endolytic microbial community of these rocks to directly demonstrate the biogenesis of the

moonmilk. Here we present an example of an ongoing co-evolution between rocks and microorganisms: the moonmilk

100   contributes to the evolution of the rock rich in calcium carbonate, while the physio-chemical features of the host rock shape

the resident microbial community which induces the moonmilk deposition.

2 Materials and Methods

2.1 Site description and sampling

105 Samples representative of the bedrock were collected from the ancient Etruscan Necropolis of Tarquinia, a UNESCO World Heritage site (Viterbo, Italy), in which more than 200 painted hypogeal tombs (dated from the 7th to the 2nd century B.C.) were discovered. The tombs were excavated in a sedimentary bedrock belonging to a middle to upper Pliocene formation known as *Macco* consisting of yellowish bioclastic calcarenites interbedded with hybrid sandstones (Supplementary Fig 1. shows the map of the Necropolis in Tarquinia and the sampling locations: Macco s.s. (red dots) and Hybrid sandstone (blue

110 dots). From each location, rock samples were collected and kept in plastic bags on ice and transported to the laboratory for analyses. With sterile hammer and chisel, surface material, samples from outdoor and indoor rocks were first removed to a depth of 3 to 5 cm. The interior of the rock samples was processed for DNA extraction or geological experiments.

2.2 DNA extraction procedures and sequencing of rock and moonmilk samples

For each sample, rock material was collected aseptically using a sterile rock hammer or chisel and stored in sterile collection

115 bags. All samples intended for DNA extractions were collected by discarding the top 3-5 cm layer and then crushing the inner part with a sterile rock hammer and further reducing it to a powder by grinding with a sterile mortar and pestle. Genomic DNA extraction was performed using the DNeasy PowerMax Soil Kit (QIAGEN) following the manufactor manual method using about 10g of collected material. Spectrophotometric quantification was performed using a ThermoScientific NanoDrop spectrophotometer (Thermoscientific), and DNA purity was assessed through the evaluation of 260/280 and 260/230

120 absorbance ratios. PCR amplification was performed on about 50 ng of DNA of each sample as described by Grottoli et al., (2020). Amplicon sequencing was performed on PCR products deriving from the 16S rRNA samples regions. PCR products were sequenced through Oxford Nanopore Technologies (ONT) by a MinION sequencer. A total amount of 200 ng of PCR products for samples were used for sequencing. Rapid barcoding of PCRs was carried out following the protocol released by ONT (SQK-RBK004) and sequenced by a Flongle flocell (FLO-FLG001). Total run produced ~15 Mb of data (~34 k reads)

125 including: ~2.8 Mb (~4 k reads) for Moonmilk of Tomba Maggi 2, ~2.8 Mb (~4 k reads) for Moonmilk of Tomba degli Scudi, ~2.7 Mb (~11 k reads) for hybrid sandstone and ~6.4 Mb (~15 k reads) for calcarenite, respectively. Kraken2 system (Wood et al., 2019) produced a taxonomic classification of 98.83%, 96.65%, 44.02% and 62.29% of total reads for each sample, Moonmilk of Tomba Maggi 2, Moonmilk of Tomba degli Scudi, hybrid sandstone and calcarenite, respectively.

2.3 Petrophysical analysis

130 Polarized light microscopy (PLM) observations of rock samples and moonmilk speleothems have been carried out on thin-sections at the Dipartimento di Scienze della Terra of Sapienza University of Rome by means of Olympus BX50 (Japan) and

Zeiss Axiophot (Germany) transmitted polarized light microscopes, under plane-polarized light and crossed polars, at 25×, 40×, 100 ×, 200 ×, 400 × magnifications. Microscopic images were obtained by using a digital camera. Polished thin sections (28 mm x 48mm) were prepared from the samples vacuum impregnated with epoxy resin before cutting and thinning. Density and porosity were measured using Ultra-pyc 5000 helium pycnometer from Anton Paar with an accuracy of 0.02% and repeatability of 0.01%. Bulk density was obtained by dividing the dry mass of sample by its total volume. Grain density results from the calculation of the mass / measured volume ratio of the pulverised matrix. Both total and effective (open) porosity was measured. The latter was obtained by dividing the difference between the geometric volume and the volume measured by pycnometer of the sample (Ruggieri and Trippetta, 2020; Trippetta et al., 2020). All laboratory measurements were made in the Earthquake Physics Laboratory at Sapienza Earth Sciences Department. Bulk rock major and trace element compositions were obtained by lithium metaborate/tetraborate fusion ICP-AES and ICP–MS, at Activation Laboratories Inc. (Ontario, Canada) according to the Code 4Litho code package on solutions prepared with lithium metaborate fusion. Loss on Ignition (LOI) was measured according to standard gravimetric procedures. Details on the precision and accuracy of the analyses are reported in www.actlabs.com.

The calcium carbonate content was assessed by gasometric measurements using a Dietrich–Fruhling calcimeter measuring one gram of the bulk sediment following Siesser et al., (1971).

2.4 Scanning Electron microscopy analysis

Scanning Electron Microscopy (SEM) was performed on the Moonmilk samples and on the rock thin sections using a Field Emission Scanning Electron Microscopy (FESEM) Zeiss Auriga 405, with a chamber room that maintains a pressure of about 10^{-5} to 10^{-6} mbar. Before mounting the samples inside the microscope, the specimens were coated with 20 nm of chromium using a Quorum Q150T sputter. Chromium has a high X-ray Kα value (5.145 keV), so does not interfere with lighter elements during the EDX analysis. EDX spectra were obtained using a Bruker Quantax detector in point mode for 30 s, with the electron microscope acceleration voltage set at 10 kV and working distance of 6 mm to optimize the number of the incoming X-ray signal.

3 Results and Discussion

3.1 Lithology, mineralogy and geochemical characterization of the bedrock

The area of the Monterozzi Necropolis offers a favourable chance to investigate the role of bedrock in the genesis of the moonmilk. To this purpose, we analysed samples of bedrock taken from different areas of the Necropolis with special focus to the sites on which *Tomba dei Vasi dipinti*, *Tomba Maggi 2, Tomba delle Pantere* and *Tomba degli Scudi* were carved (Supplementary Fig. 1). The hypogeal tombs of the Monterozzi Necropolis are excavated within a sedimentary substrate of Middle to Upper Pliocene age known as *Macco* formation consisting of two main lithofacies showing lateral and vertical heteropic relationships. *Macco* s.s. lithofacies is a bioclastic calcarenite represented by packstone to rudstone and floatstone

composed of small volume (<5-10%) of micrite matrix, coralline algal branches, bryozoans, bivalves (pectinids and oysters),
echinoids, benthic foraminifers, and skeletal debris. Rare non-carbonate grains are present. In the inner walls of the
intergranular voids microsparite cement precipitation and/or recrystallization often occurs, due to diagenetic processes of
dissolution. The second lithofacies is represented by a poorly cemented, crudely stratified, hybrid sandstone. It is fine to
medium grain-sized and grain-supported, with small amount of micrite matrix (< 10% vol) and nearly devoid of carbonate
cement. This lithofacies is characterised by abundant bioturbation, the skeletal assemblage is dominated by small benthic
foraminifers, echinoids, serpulids and bivalves (mainly oysters). Planktonic foraminifers are common. The terrigenous fraction
mainly consists of monocrystalline grains of quartz, sedimentary lithoclasts, and subordinate detrital micas and feldspar along
with rare glaucony grains and opaques. Calcimetric analyses from the two lithofacies revealed that calcium carbonate contents
range from 90 to 98% in the Macco s.s. calcarenite and from 49 to 59 % in the hybrid sandstone. Moreover, whole rock major
oxide and trace element compositions highlight the geochemical difference between the two lithofacies (Table 1), mostly
related to the higher proportion of terrigenous fraction in the hybrid sandstone. Helium pycnometry revealed a high open
porosity in both lithofacies (~43% for Macco s.s.; ~42% for hybrid sandstone). The *Macco* s.s. lithofacies shows a dominance
of vuggy porosity, abundant intraparticle, interparticle and mouldic porosity. The main porosity of hybrid sandstone is
represented by interparticle porosity, rarely by mouldic porosity. Thus, despite the differences in composition, the calcarenite
and the hybrid sandstone show two essential characteristics required for moonmilk formation: high calcium content, which
activates the microbial metabolism leading to biomineralization, and high porosity, necessary for the exchange of fluids and
nutrients in an oligotrophic environment, also providing the space for microbial colonization.

3.2 First report of inner location of moonmilk

Although with some differences, moonmilk biogenic deposits develop indifferently within the two distinct lithofacies
constituting the bedrock of the Etruscan necropolis of Monterozzi (Mura et al., 2021). Fig. 1a shows an example of hypogeal
walls of tombs carved in calcarenite and hybrid sandstone, and the walls covered by moonmilk in the *Tomba degli Scudi* and
Tomba Maggi 2 (Fig. 1b). The moonmilk layer originated from hybrid sandstone is thinner than the one observed on a
calcarenite substrate, but the scanning electron micrographs of the moonmilk sampled from the walls of *Tomba degli Scudi*
and *Tomba Maggi 2* showed the same nanofiber structure (Fig. 1c). X-ray powder diffraction analysis (XRD) revealed that
moonmilk is composed of calcite (Mura et al., 2020).

By now, the moonmilk has only been considered as a deposition covering rock surfaces (Borsato et al., 2000), but the analysis
with transmitted polarized light microscopy of thin sections showed that the moonmilk is present *inside* the calcarenite in the
vuggy and moldic porosity of the calcarenite in *Tomba Maggi* 2 (Fig. 2a) and in intergranular and moldic pores of the hybrid
sandstone bedrock collected inside the *Tomba degli Scudi* (Fig. 2b). The presence of moonmilk inside the rocks is a general

phenomenon, because it is observed in all samples, regardless the type of rock, calcarenite or hybrid sandstone, and where
195   were collected, outdoor or indoor (Supplementary Fig. 2, 3 and 4).

These results are also supported by the discovery of moonmilk deep inside a calcarenite rock sampled at the entrance of the
Tomba dei Vasi Dipinti (Supplementary Fig. 5).

The analysis of the rock substrate sampled outdoor the *Tomba dei Vasi Dipinti* also suggests that the moonmilk could contribute
to the authigenic carbonate growth in the host rock, covering the inner walls of the voids (Fig. 3).

200   **3.3 Co-evolution of rocks and microorganisms**

If the moonmilk that was observed *inside* the rocks (a location that was not reported before) is of biogenic origin, traces of
biomineralization would be expected. Indeed, the SEM analysis on the thin sections of calcarenite sampled outdoor of the
Tomba dei Vasi Dipinti revealed many structures corresponding to biomineralization (Fig. 4 and Supplementary Fig. 6).

Bacterial biomineralization was also detected in the sandstone sampled indoor the *Tomba degli Scudi* (Supplementary Fig. 7)
205   and in calcarenite sampled outdoor of the *Tomba delle Pantere* and indoor of *Tomba Maggi 2* (Supplementary Fig. 8),
suggesting that this is a common phenomenon. Such bacterial biomineralization is also known as "*entombment*" and it is easily
observed in laboratory settings when bacterial strains are subjected to environmental conditions favoring calcium carbonate
precipitation (Supplementary Fig. 9a, b, c). In plates, the precipitation of calcium carbonate occurs even at a considerable
distance from the bacterial colony, possibly by the diffusion of extracellular enzymes known to be involved in calcium
210   carbonate metabolism (Dhami et al., 2014; Rodriguez-Navarro et al., 2019) (Supplementary Fig. 9d). It remains unclear how
the moonmilk nanofibers are produced in natural environmental conditions because to date it has been impossible to reproduce
its formation in the laboratory. In fact, bacterial strains cultured from rocks represent only a negligible fraction of the total
microorganisms present in the rocks. Instead, the entire microbial community, with a metabolism that sustains the growth in
the rock environment, is needed to precipitate and dissolve calcium carbonate.

215   Overall, our results underscore the role of microorganisms in promoting moonmilk deposition, contributing to the rock
formation processes. Nevertheless, to propose the moonmilk as part of a geological process, the microbial communities of the
rocks and those contributing to the moonmilk deposition should have a similar composition. Aiming to identify the rock
microbial communities, samples from calcarenite and hybrid sandstone were analysed together with the corresponding
moonmilk samples from *Tomba Maggi 2* and *Tomba degli Scudi*. The results of the 16S rRNA amplicon sequencing showed
220   a high abundance of Actinobacteria, Bacteroidetes, Cyanobacteria, Firmicutes and Proteobacteria (Fig. 5).

Bacterial community diversity was measured by inverse Simpson index and Shannon index for moonmilk (*Tomba degli Scudi*
and *Tomba Maggi 2*) and rocks (calcarenite and hybrid sandstone). The indices do not show any significant differences between
the samples (Mann Whitney test, P > 0.05) (Supplementary Fig. 10). These results show that in moonmilk and rocks the
microbial composition is similar, irrespective of rock type (calcarenite and hybrid sandstone) or the environment where the
225   samples were collected (outdoor or indoor).

It should be noted that 16S RNA analyses do not provide information about metabolic activity, thus these data do not identify microorganisms that are active in $CaCO_3$ deposition, but the overall data demonstrate that it is the endolytic community of the rocks is promoting moonmilk deposition; the results presented also revealed the presence of biomineralization and calcite nanofibers that originate from bacterial entombment, not only on the surface, but inside the rocks, demonstrating the biogenic

230   origin of the moonmilk. The presence of a resident microbial community deep within the rocks, possibly evolved with the rocks through geological time. Therefore, no habitat should be considered as *extreme* for the resident microbial community, and the rocks should not be considered as a 'refuge' for escaping extreme environmental conditions. Biological research should focus on microbial community evolution with respect to the geologic substrate in which they are living, considering the natural co-evolution of microbes and rocks, and possibly abstaining from the consideration of the microbial metabolism as an

235   adaptation to *adverse* environmental conditions.

3.4 Moonmilk as a biosignature

The search for traces of extra-terrestrial life is a complex task often ending with inconclusive results. The co-evolution of minerals and microorganisms has implications for the quest of evidence for life on other planets. The discovery of minerals of undisputable biological origin, rather than organic remains, may provide the most robust signs of biological activity (Hazen et

240   al., 2008). Coevolution of life and minerals throughout Earth history lays the ground for an inclusive search for the presence of life, because not only rocks arisen from life but also because life itself may have formed from rocks (Bizzarri et al., 2021; Marshall, 2020; Saladino et al., 2018). Thousands of earth's minerals owe their existence to the development of life on the planet and calcium carbonate phases that are massively produced on earth by microorganisms, are the best example (Hazen et al., 2008).

245   We have focused on the calcium carbonate speleothems called moonmilk. Given the tight association and co-evolution between rocks and microbial communities that results in the observed biomineralization, calcite nanofibers are of interest in the field of astrobiology and are considered as a potential sign of life.

The moonmilk production contributed to rock formation by filling the pores and the cracks in the rocks, while the rock composition and the porosity shaped a microbial community that copes with high calcium content producing calcite nanofibers.

250   The moonmilk is mainly found in karst caves, but there are examples of moonmilk bioprecipitation also in hypogeal environments carved in different geologic substrates, such as granitoid rocks or sandstone (Saladino et al., 2018; Miller et al., 2018). The moonmilk has been also found in lava tubes where the microbial communities are similar to those present in the moonmilk that originated from calcarenite (Miller et al., 2020; Gonzalez-Pimentel et al., 2021), raising the possibility of extending the moonmilk as a biosignature also beyond the Earth calcium carbonate rocks.

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260 *Author contributions.* S.R., M.B., A.C. and F.M. performed the experiments; M.B., S.R., T.R. collected samples; R.N., A.G. and M.R. supervised and performed the nucleic acid analysis; D.F.M supervised the sampling in the Etruscan tombs; T.R. and E.D.M. conceived the study and T.R. wrote the paper with contributions from S.R., M.B and E.D.M.

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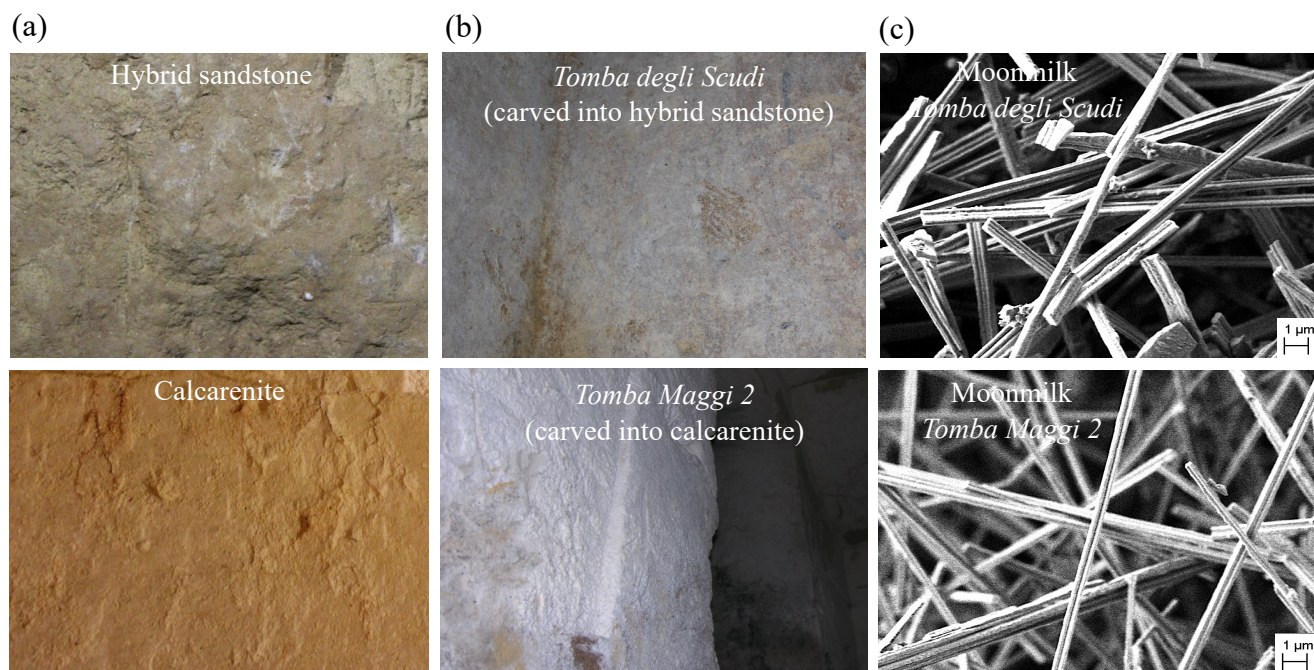
Figure 1. In Tarquinia, Italy, during the Iron Age, the ancient Etruscans carved hypogeal tombs into calcarenite and in hybrid sandstone bedrock whose walls are covered of the moonmilk, a secondary speleothem. (a): examples of hypogeal walls of tombs carved in hybrid sandstone and calcarenite, the absence of the moonmilk is due to the restoration interventions. (b): Along the centuries, the moonmilk speleothem precipitated as a white patina on the walls and ceilings of the tombs: as an example, the walls covered of moonmilk of the *Tomba degli Scudi* and the *Tomba Maggi 2* are shown, carved in hybrid sandstone and calcarenite, respectively. (c) Scanning electron micrographs of the moonmilk sampled on the walls showed in (b) in the *Tomba degli Scudi* and the *Tomba Maggi 2*. Regardless the rock substrate in which the moonmilk is formed, the structure of the nanofibers are similar.
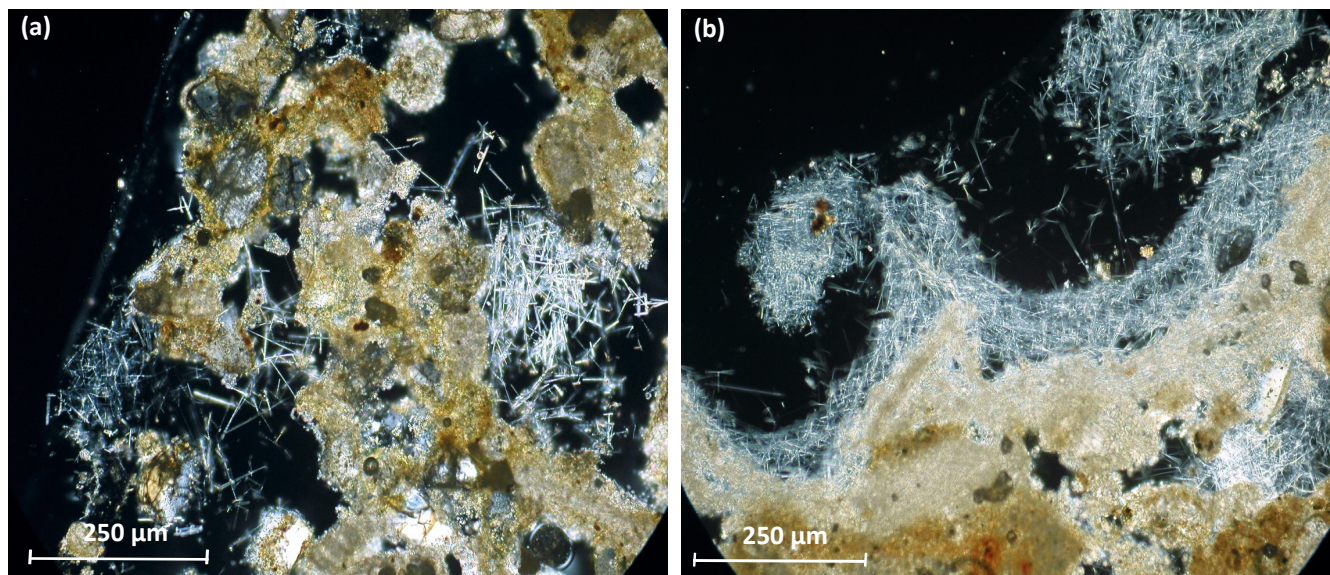
Figure 2. The moonmilk is present on the surface and inside the calcarenite and sandstone rocks.
Optical microscope thin-section micrographs in crossed-polarized transmitted light. (a): calcarenite sample collected inside the *Tomba Maggi 2; (*b): hybrid sandstone sample collected inside the *Tomba degli Scudi.*

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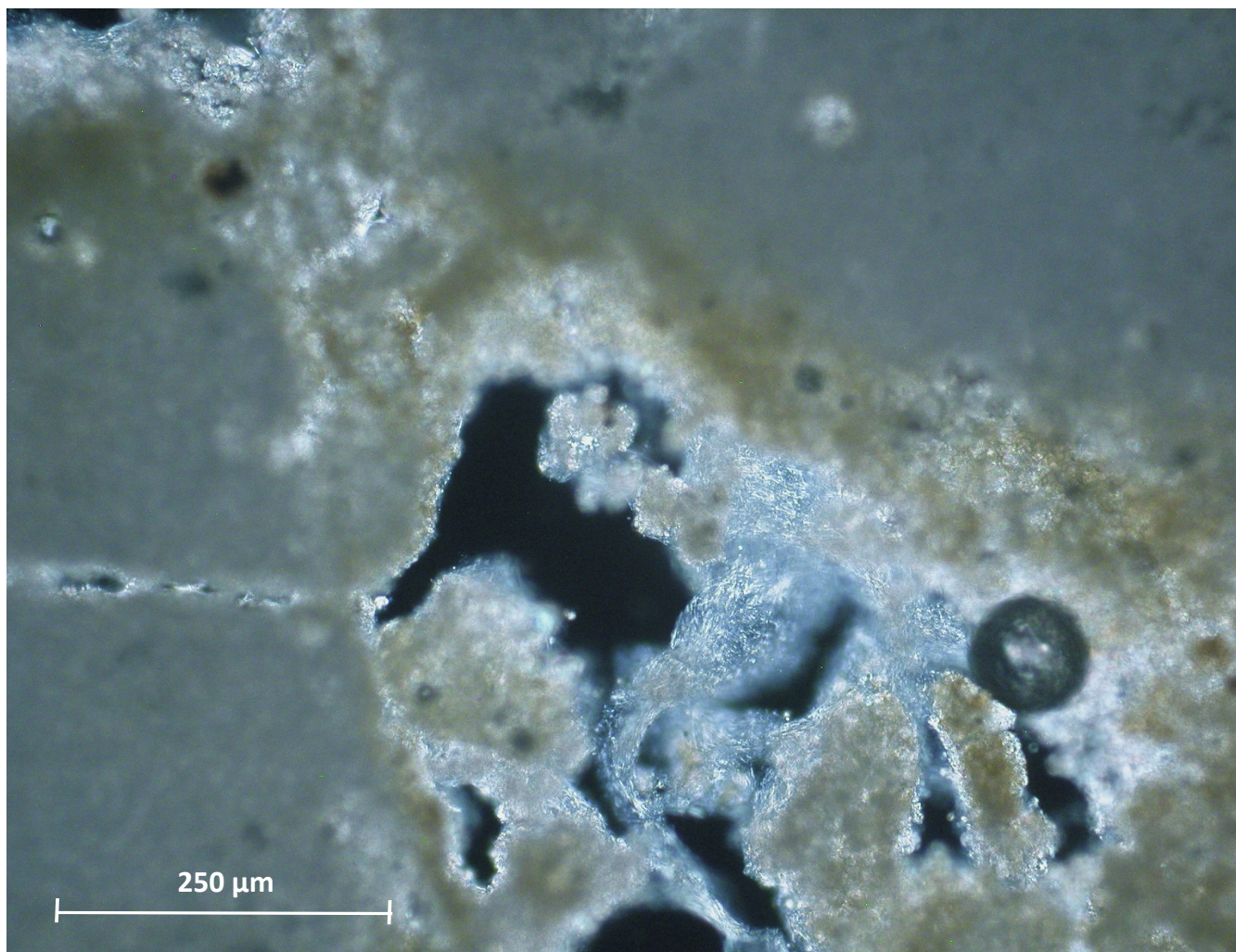
Figure 3. The moonmilk contributes to the lithogenic processes. Optical microscope thin-section micrographs (crossed polarized transmitted light) of the moonmilk speleothems grown into the calcarenite sampled outdoor of *Tomba dei Vasi Dipinti*.
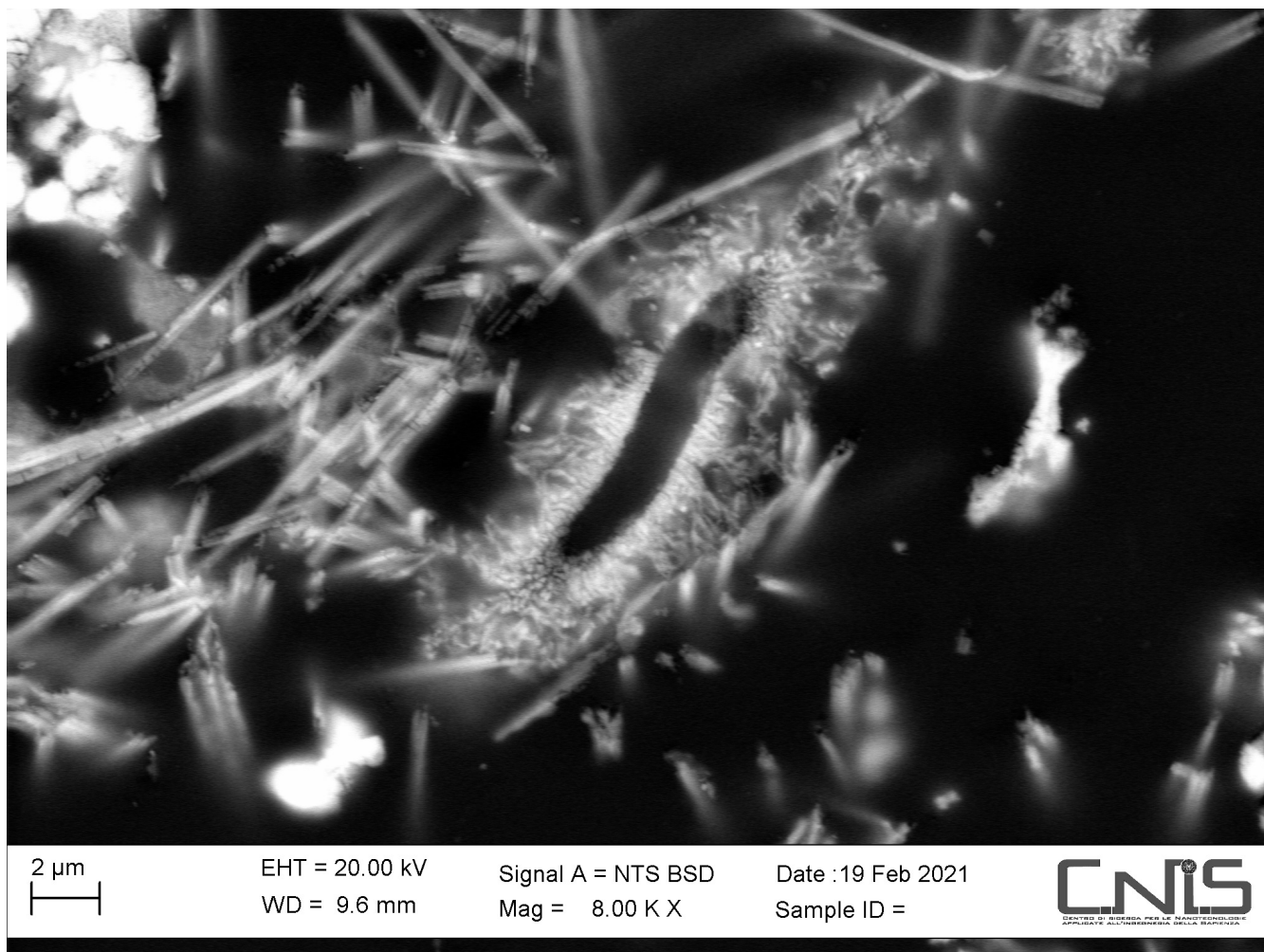
Figure 4. Bacterial biomineralization in the calcarenite. *S*canning electron micrograph of a thin-section of the calcarenite sampled outdoor of the *Tomba dei Vasi Dipinti.*
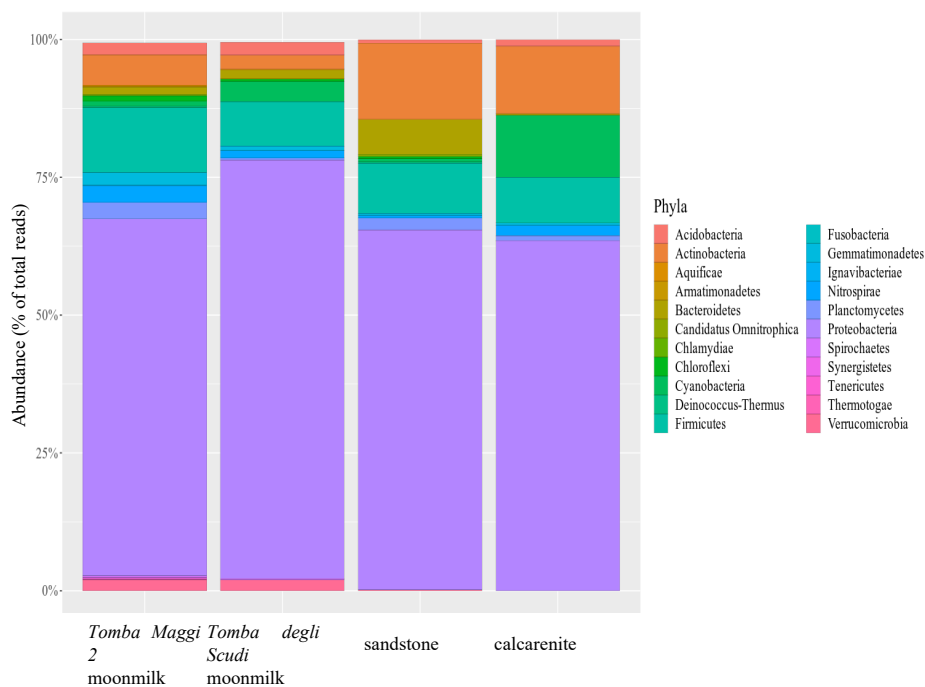
Figure 5. Phyla present in the microbial community from moonmilk samples of *Tomba Maggi 2, Tomba degli Scudi and* their corresponding rocks (calcarenite and hybrid sandstone). The histogram shows the Phylum relative abundance (%) for the analysed samples. Community structure was determined by targeted amplicon sequencing of bacterial 16S rRNA genes. All samples show a high abundance of Actinobacteria, Bacteroidetes, Cyanobacteria, Firmicutes and Proteobacteria.