SILICICLASTIC MICROSTROMATOLITES IN A SANDSTONE CAVE: ROLE OF TRAPPING AND BINDING OF DETRITAL PARTICLES IN FORMATION OF CAVE DEPOSITS

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Abstract: The article deals with finely laminated microstromatolites composed of detrital siliciclastic particles, mainly quartz, feldspars and clay minerals, lining the walls of W Sopotni Wielkiej Cave (Polish Outer Carpathians). Newly precipitated mineral phases do not contribute to their growth. The microstromatolites cover vertical and overhanging walls of the cave. They form subhorizontal ripples and tongue-shaped stepped microrracettes. The stromatolites are constructed by bacteria and Actinomycetes. Seven morphotypes of micro-organisms have been distinguished. Trapping and binding of detrital particles result in the microstromatolite growth. The growth is influenced by the relatively close distance to the soil cover which provides detrital mineral particles and by the presence of gravitationally widened fissures which guide the water transporting mineral particles down to the cave. The particles are transported only during wet periods. The episodic supply of the particles results in visible lamination of microstromatolites. The microrracettes form in zones of increased water-flow. The lack of autochthonous components most probably reflects too low saturation of water to precipitate any minerals.

Key words: bacteria, Actinomycetes, biofilm, speleothems, Outer Carpathians.

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INTRODUCTION

Constructive role of micro-organisms in speleothem formation is a focus of growing interest (see review by Northup & Lavoie, 2001; Jones, 2001, 2010; Taboroší, 2006 and references quoted therein). A great body of literature exists on the importance of bacteria and fungi for precipitation of carbonate speleothems, especially for the growth of moonmilk, cave pisoids and some subaqueous formation (e.g., Gradziński et al., 1997b; Gradziński, 2000; Melim et al., 2001; Jones, 2009a; Curry et al., 2009). The plausible direct or indirect influence of these micro-organisms on the crystallization of needle-fibre calcite in cavern environment is commonly discussed (Blyth & Frisia, 2008; Richter et al., 2008). The precipitation of speleothem-forming manganese and iron oxides, opal and phosphates is also mediated by biological activity (e.g., Gradziński et al., 1995; Manolache & Onac, 2000; Aubrecht et al., 2008; Jones, 2009b).

Conversely, little attention has been drawn to trapping and binding of detrital grains of various mineral composition as a constructive mechanism in speleothem growth. Such mechanism has been mentioned on the occasion of studying the microbially driven precipitation of diverse speleothems in cavern environment. However, it has only been regarded as a subordinate contribution to speleothem growth. Jones and Motyka (1987) showed that algae trap and bind carbonate micrite, which eventually leads to formation of the well visible lamination. Cyanobacteria, representing Gleocapsa sp., are capable of trapping air-born quartz grains and insect fragments, which collectively contribute to the formation of crayfish-like profile stalagmites located in entrance parts of Australian caves (Cox et al., 1989a, b; James et al., 1994). Jones (1995) noted some examples of trapping and binding of detrital grains to the sub-strate by organic filaments in the biofilm covering walls in twilight zones of caves. Cunningham et al. (1995, fig. 7) illustrated binding of corrosion residuum by filamentous micro-organisms in Lechugilla Cave (New Mexico). Cañaveras et al. (2001) mentioned trapping and binding of
micritic grains resulting from substrate breakdown in the famous Altamira cave. Gradziński and Holubek (2005) found detrital silicate and dolomite grains incorporated into subaqueous cottonball speleothem in Zlomiská Cave (Slovakia); the cottonballs presumably originated under biological mediation. Baskar et al. (2006) suggested that detrital grains organically trapped and bound are incorporated into carbonate speleothems in Borra Caves (India). Also, biologically mediated opal speleothems from Venezuelan caves contain some grains trapped and bound by micro-organisms (Aubrecht et al., 2008). Detrital grains contribute greatly to formation of some speleothems in sandstone caves in the Polish Outer Carpathians (Urban et al., 2007). In some cases they compose deposits whose relief imitates the morphology taken by speleothems. Some of these forms are cemented with opal.

Clastic formations built of sand- and silt-sized grains are present in many caves (Hill & Forti, 1997, p. 219–221 and references therein). Some of them are akin to carbonate speleothems in their shape and dimension although they are only cemented by carbonate minerals (e.g., Baker, 1942). In fact, many of them represent erosional remnants after accumulation of cave clastics (e.g., Gradziński & Radomski, 1960) and they do not share their origin with typical speleothems. Vermiculations represent another type of cave formation composed of siliciclastic material (Hill & Forti, 1997, p. 221–223). They are found on cave walls or ceilings as irregular spots or bifurcating stripes. Their array gives the overall impression of a group of warms or a tiger or leopard skin. In the majority of papers no suggestions have been made on the possible role of micro-organisms in the origin of the above clastic cave formations, excluding special type of vermication in sulfidic caves. Although Bini et al. (1978) considered influence of bacteria on formation of vermication, ultimately they ruled out this influence.

This paper documents the presence of microstromatolites composed of detrital siliciclastic particles lining the walls of a sandstone cave. The significance of micro-organisms to trapping and binding detrital particles, and therefore to forming the microstromatolites in question, is discussed. A hypothesis is put forward on a similar constructive role of micro-organisms in formation of clastic-rich laminae within carbonate speleothems.

**GEOLOGICAL AND SPELEOLOGICAL SETTING**

Siliciclastic microstromatolites were found in W Sopotni Wielkiej Cave (Jaskinia w Sopotni Wielkiej) and reported during a local symposium (Gradziński et al., 2001). The cave is located in the Beskid Żywiecki Mts. From the geological point of view this region belongs to the Outer Carpathians, which are built predominantly of Cretaceous–Palaeogene flysch (Fig. 1). The cave is formed in the Eocene thick-bedded sandstones of the Magura beds belonging to the Rača Unit of the Magura Nappe (Golonka & Wójcik, 1978). The sandstone is fine grained. An X-ray diffractogram of the sandstone reveals the presence of quartz, alkali feldspars and two groups of phyllosilicates, with a major order layer separation of 14.5 Å and 10.0 Å (see Figs 4A, 5A). The 14.5 Å minerals are either chlorite or smectite, although further analyses are necessary for exact identification. The 10.0 Å peak may indicate the presence of illite or micas. Observations in optical microscope revealed the presence of muscovite and biotite micas (see Fig. 5A); a confirmation of the illite presence requires < 2 µm fraction separation.

The cave main entrance is located at the altitude of 840 m (Klassek, 1997). The cave is composed of a series of passages up to 2 m wide and 5 m high, with total 101 m in length (Fig. 2). The passages display rectangular or triangular cross-sections. Their floors are littered with sandstone debris. The cave originated due to widening of fractures under the action of downslope creep of sandstone blocks towards the neighbouring valley. Hence, it represents a crevice type cave (sensu Palmer, 2007, p. 7). The cave is protected as a ‘nature monument’.

The microstromatolites cover the walls of the Komora Trójkątna (Triangular Chamber) located ca. 12 m from the cave entrance. They were noted by Mikuszewski (1973) and Klassek (1997) but recognized as carbonate flowstones. The chamber is completely dark; its walls are covered by drops of condensation water. The internal temperature is 5.5 °C (Klassek, 1997). The thickness of the roof rock over the Triangular Chamber is estimated at 6–8 m. The area over the cave is forested.

**MATERIAL AND METHODS**

**Morphology and internal structures**

Microstromatolites were documented in the Triangular Chamber (Fig. 2). Sampling was limited to the minimum due to the cave protection. Samples were collected from
both sides of the chamber, preferably from hidden places. A thin section was made from one sample sunken in a box filled with ARALDITE. Internal structures of the microstromatolites were observed under petrographic and scanning electron microscope (SEM) JEOL 5410, coupled with a microprobe (EDS) Voyager 3100 (Noran product). The samples were mounted on SEM holders with silver glue and coated with C or Au. Samples were lyophilized prior to coating to prevent collapse of the organic structure.

Mineral composition

Samples were analysed by the powder X-ray diffractionmetry (XRD) using a vertical XPer APD Philips goniometer (PW 1830). Infrared absorption spectra (IR-FT) were obtained at ambient temperature and with 2.0 cm⁻¹ resolution using a BIO-RAD Fourier Transform Spectrometer (FTS 135).

Microbiology

Samples were aseptically collected to sterile glass flasks, transported to laboratory and suspended in physiological salt solution, shaken and inoculated in liquid and agar media. They were incubated at 20 °C and 35 °C from 1 to 21 days. The growth of micro-organisms was systematically monitored. The following microbiological media were used for isolation: Beef Extract − Nutrient Broth − Merck, Trypticase Soy Broth (Soybean-Casein Digest Medium) − BioMerieux, Nutrient Agar − Merck, TSA (Tryp case Soy Agar) − BioMerieux, Soil Extract Agar (Atlas & Parks, 1997), Iron Bacteria Isolation Medium (Atlas & Parks, 1997) and Actinomycetes Isolation Agar (Atlas & Parks, 1997). After incubation the clean cultures of bacteria were isolated on agar media (Pepper & Gerba, 2004). Morphology, Gram stain and biochemical proprieties of bacteria were analyzed to identify micro-organisms. Species identification was based on Bergey’s Manual of Determinative Bacteriology and Bergey’s Manual of Systematic Bacteriology (Holt, 1989, 1994). Since there are no standard biochemical tests for the majority of isolated genera, the biochemical tests were individually selected according to diagnostic manuals.

RESULTS

Relief

The microstromatolites display a soft, pasty consistency and contain a substantial amount of water. They cover the vertical and overhanging walls of the Triangular Chamber (Fig. 3A, B, D). The dip of rocky walls covered with the microstromatolites ranges from 90° (vertical) to 110° (overhanging). The microstromatolites occur also on vertically oriented convex bends of a cave wall, which overhangs with an angle between 93° and 120° (Fig. 3A, B). Recently, they coat around 0.5 m² of the southern wall of the chamber (A in Fig. 2) and around 1 m² of its opposite wall (B in Fig. 2). However, it is very plausible that they formerly occupied a much bigger area and have been destroyed by visitors. On the vertical walls, they form more or less horizontal ripples with relief (distance between the wall and the crest), up to

Fig. 2. Simplified map of W Sopotni Wielkiej Cave (after Klassek, 1997), sampling sites are indicated (white arrows), big black arrow shows the cave entrance
2 mm (Fig. 3D). The length of particular ripples exceeds 35 cm. The neighboring ripples bifurcate. With the wall dip changing towards more overhanging, the ripples become more sinuous and form tongue-shaped stepped microterracettes (Fig. 3A, B, D). The microterracettes overhang and their upper parts dip outward at a maximum angle of 45°. In some cases, a microrim is developed along their crest with a micropool formed behind it. Microterracettes are particularly well developed along a vertically oriented convex bends of the cave wall (Fig. 3B). The distance between the microterracette crest and rock wall reaches 1.5 cm. The vertical distance between neighbouring ripples ranges between 2 and 5 mm, whereas between microterracettes it is slightly higher and maximally reaches 8 mm.

**Microbiology**

Microbiological analysis reveals the occurrence of several taxa which are listed in Table 1. It seems reasonable to accept that Arthrobacter, Bacilli and Actinomycetes belong to indigenous microflora of the analysed microstromatolites. A scarce presence of Micrococcus and Staphylococcus in the studied samples implies that they should be considered allochthonous. A preferred environment of their growth supports the above view (Table 1).

**Mineral composition**

Quartz is the major constituent of the microstromatolites; the alkali feldspars and clay minerals have a smaller
contribution. The clay mineral content, although insignificant, is slightly higher in the microstromatolites than in the host rock. Sheet silicates show diffraction peaks at 14.5  and 10.0  as well as a minor peak centered at 12.0  (Fig. 4A). The 14.5  and 10.0  peaks correspond probably to minerals present in the host rocks (smectite or chlorite and micas), whereas the 12.0  peak reflects most likely the presence of mixed-layer clay minerals. This issue requires further investigation.

Additional information as to the composition of the microstromatolite sample is brought by IR absorption spectrum (Fig. 4B). A weak absorption band at 1405 cm−1 may point to the presence of only very small amounts of carbonates. The weak intensity of this absorption band and the absence of appropriate diffraction peaks, that is 3.03  (Fig. 4A) indicate almost negligible amount of carbonates. The microstromatolites have similar composition to the sandstone which hosts the cave (Fig. 5A). The XRD patterns are given for comparison of the mineralogy of both rocks (Fig. 4A).

Internal structures

The surface of the microstromatolites is uneven. It displays elevated clumps surrounded by depressions. The clumps show lower porosity than the depressions. Also the microrim crest is characterized by a lower porosity than the micropool fringed by it (Fig. 5B).

The microstromatolite is finely laminated, with the lamina thickness ca. 50–200 µm visible due to differentiation in mineral composition, grain size, and probably changes in microporosity (Fig. 5C). The laminae are convex outward, irregular. Some of them have confused boundaries. They are composed of particles of clay and fine silt fraction. Sporadic quartz and muscovite grains reach up to 150 µm across (Fig. 5C). Outsized quartz and muscovite grains are concentrated in thicker laminae.

### Table 1

<table>
<thead>
<tr>
<th>Bacteria</th>
<th>Occurrence</th>
<th>Morphology</th>
<th>Properties</th>
</tr>
</thead>
<tbody>
<tr>
<td>Arthrobacter sp.</td>
<td>sample 1, 2</td>
<td>Gram positive irregular, nonsporing rods, in old cultures cocci</td>
<td>Aerobic, chemoorganotrophic; grows on simple media, widely distributed, principally in soils; psychrothropic arthrobacters have been reported to predominate in subterranean cave silts</td>
</tr>
<tr>
<td>Amycolata autotrophica</td>
<td>sample 1, 2</td>
<td>Gram positive, Actinomycetes, branched vegetative hyphae</td>
<td>Aerobic, chemoorganotrophic - facultatively autotrophic; isolated from diverse habitats</td>
</tr>
<tr>
<td>Bacillus alcalophilus</td>
<td>sample 1</td>
<td>Endospore forming, gram positive rods</td>
<td>Aerobic, chemoorganotrophic; alcali tolerant, isolated from various material in media at pH 10</td>
</tr>
<tr>
<td>Bacillus megaterium</td>
<td>sample 1, 2</td>
<td>Endospore forming, gram positive rods, cell diameter over 1 µm</td>
<td>Aerobic, chemoorganotrophic; found in wide range of habitats; grows in low temperatures</td>
</tr>
<tr>
<td>Bacillus mycoides</td>
<td>sample 1</td>
<td>Endospore forming, gram positive rods, cell diameter over 1 µm, forms filaments</td>
<td>Aerobic but can grow anaerobically, chemoorganotrophic; forms rhizoid colonies; widely distributed</td>
</tr>
<tr>
<td>Micrococcus varians</td>
<td>sample 2</td>
<td>Gram positive cocci, nonsporing</td>
<td>Aerobic; micrococci are common on mammalian skin, soil, air</td>
</tr>
<tr>
<td>Staphylococcus xylosus</td>
<td>sample 1</td>
<td>Gram positive cocci, nonsporing</td>
<td>Facultatively anaerobic; staphylococci are mainly associated with skin, but often isolated from food products, dust and water</td>
</tr>
<tr>
<td>Streptomyces sp.</td>
<td>sample 2</td>
<td>Gram positive, Actinomycetes, extensively branched vegetative hyphae</td>
<td>Aerobic; chemoorganotrophic; widely distributed and abundant in soil</td>
</tr>
</tbody>
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Fig. 4. A – XRD pattern of host rock and microstromatolite; Q – quartz, Al – albite, Kf – K feldspar, C14 – clay minerals with main reflection 14.5 ; C12 – clay minerals with main reflection 12.5 , C10 – 10.0  micas or illite, B – IR absorption spectrum of the microstromatolite
Observation under SEM reveals the internal structures of the microstromatolites. They are composed of mineral grains, micro-organisms and their extracellular polymeric secretions (EPS).

Clay minerals compose clumpy aggregates, up to 200 µm across (Figs 5B, D, 6A). The aggregates also comprise some admixtures of bigger mineral grains. Mica grains display angular shape and tabular habit (Fig. 6B). They contain Al,

Table 2

<table>
<thead>
<tr>
<th>Morphotype</th>
<th>Shape</th>
<th>Dimensions</th>
<th>Branching</th>
<th>Morphology</th>
<th>Cell arrangement</th>
<th>Comments</th>
<th>Figure</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>coccoid to short rods</td>
<td>Ø – 0.8 µm length &lt; 1.5 µm</td>
<td>no</td>
<td>smooth</td>
<td>chains or three-dimensional colonies covered with EPS</td>
<td>co-occur with no 6</td>
<td>6C, 7A-D</td>
</tr>
<tr>
<td>2</td>
<td>rods</td>
<td>Ø – 0.8 µm length 2-3 µm</td>
<td>no</td>
<td>smooth</td>
<td>straight or zig-zag chains</td>
<td></td>
<td>7E</td>
</tr>
<tr>
<td>3</td>
<td>spindle-shaped rods</td>
<td>Ø – 1 µm length 4-5 µm</td>
<td>no</td>
<td>granulated</td>
<td>chains</td>
<td></td>
<td>7F</td>
</tr>
<tr>
<td>4</td>
<td>coccoid</td>
<td>Ø – 0.5–0.8 µm</td>
<td>no</td>
<td>smooth</td>
<td>single or in linked pairs</td>
<td></td>
<td>8D</td>
</tr>
<tr>
<td>5</td>
<td>irregular ovoid</td>
<td>Ø – 6 µm</td>
<td>no</td>
<td>irregular but smooth</td>
<td>single or in linked pairs</td>
<td>co-occur with no 6</td>
<td>8C</td>
</tr>
<tr>
<td>6</td>
<td>filamentous</td>
<td>Ø – 1.1-1.3 µm length &gt; 20 µm</td>
<td>yes</td>
<td>spinose</td>
<td>filaments slightly curved</td>
<td>co-occur with no 4, 5, 7, 8A</td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>filamentous</td>
<td>Ø &lt; 0.5 µm length &gt; 15 µm</td>
<td>no</td>
<td>smooth</td>
<td>some filaments are twisted</td>
<td>co-occur with no 4, 6</td>
<td>8A-D</td>
</tr>
</tbody>
</table>
Si and K, which, along with the results of the XRD analysis, suggests muscovite (Fig. 6B). Quartz grains are more rounded and devoid of cleavage planes with Si as their major EDS detectable component. Mineral grains co-occur with, and are entombed by, micro-organisms which in many cases are tightly clung to them (Figs 6C, D, 7).

Seven morphotypes of micro-organisms have been distinguished. Their characteristics are listed in Table 2. All morphotypes are built exclusively of organic matter. Neither EDS analyses nor observation under SEM reveal any traces of cell mineralization. Micro-organisms show three-dimensional morphologies, only some of them are collapsed. The micro-organisms and their EPS occur on the surface of microstromatolites and in their shallow subsurface where they form an active biofilm. Deeper on, their amount radically decreases (Fig. 5D).

Affinity of the particular morphotypes is hard to be determined even in the light of the results of microbiological analyses. The morphotypes 1 to 3 most probably represent bacteria (Fig. 7A–C). Chains of coccoid cells, such as those of the morphotype 1, are typical two-dimensional colonies of coccoid bacteria developed due to cell divisions (Fig. 7A–C). Similarly, chains of rods characteristic for the morphotypes 2 and 3 are the effect of cell division (Fig. 7D, E). The morphotype 2 may be assigned to Arthrobacter genus, taking into account the list of determined micro-organisms and their shape and size (Table 1). Moreover, it forms a zig-zag chain of cells typical of this genus (cf. Carlile, 1979). The filamentous morphotypes 6 and 7 resemble Actinomycetes. The morphotype 6 is particularly akin to ‘hyphae morphotype 5’ described and illustrated by Jones (2009b, fig. 7A–C) from the Grand Cayman speleothems. This morphotype forms extensive three-dimensional, porous network (Fig. 8A–D). The coccoid morphotype 4 and ovoid morphotype 5 may represent both bacteria and spores of Actinomycetes (Fig. 8C, D). However, their close spatial relationship with the morphotype 6 strongly suggests the latter possibility. Although the EPS form irregularly twisted filaments or a dense layer which covers micro-organisms, none of the morphotypes is associated with a copious amount of EPS.

**DISCUSSION**

Trapping and binding of detrital particles cause the growth of the studied microstromatolites. Two mechanisms are evoked to entrap detrital grains into a stromatolite – adhesion by sticky EPS and baffling by complicated three-dimensional microbial community (Riding, 1991). In the discussed case the latter mechanism seems to be decisive, since the SEM examination of the studied samples has not re-
vealed copious amounts of EPS. Highly branched cells of Actinomycetes Amycolata and Streptomyces species act as a dense three-dimensional network capable of baffling the detrital particles. Nevertheless, Arthrobacter and Bacillus cells can excrete some slimy substances, hence the former mechanism may also work, but only to a limited extent.

Newly precipitated mineral phases seem not to contribute to the microstromatolite growth. Bearing in mind their growth mechanism, the microstromatolites in question represent agglutinated stromatolites sensu Riding (1991). In this aspect they bear a strong resemblance to some modern stromatolites in marine (Schwarz et al., 1975) and lacustrine settings (Squyres et al., 1991) as well as to several fossil examples (Martin et al., 1993; Braga & Martin, 2000 and references therein). Martin et al. (1993) coined the term siliclastic stromatolite which adequately reflects the composition and mode of growth of the studied examples. The size of entrapped siliclastic grains differentiates cave microstromatolites from the hitherto described marine and lacustrine ones. Non-spelean forms are composed mainly of sand grains with some admixtures of coarser material (Martin et al., 1993); however, some silt-rich siliclastic stromatolites are also known (Bertrand-Sarfati, 1994). Jones and Kahle (1985) introduced the term microstromatolites in order to describe microbolites, formed of fine carbonate particles and displaying fine lamination, recognized in cave sediments from the Cayman Islands. The term microstromatolite seems to be appropriate to the studied deposits. The mode of growth distinguishes the described forms from any other hitherto known microbial cave deposits being con-
constructed predominantly by minerals precipitated from solution. Although the discussed deposits mimic the relief of some speleothems (see below and Fig. 3), they cannot be included into this genetically defined group, because they contain extremely small amount of, if any, secondary mineral phases, that is minerals precipitated within a cave (cf. Hill & Forti, 1997, p. 45). The microstromatolites bear morphological resemblance to speleothems described from other sandstone caves in the Polish Outer Carpathians (Urban et al., 2007). Many of the latter forms are composed of detrital grains cemented with opal, which is not a case in the discussed microstromatolites.

The growth of the discussed microstromatolites is influenced by several factors. The influx of detrital particles is possible due to relatively close distance to the Earth surface and the weathering zone (see Urban et al., 2007). It is additionally facilitated by the presence of gravitationally widened fissures which guide the water transporting mineral particles down to the cave. The fine-grained nature of microstromatolites more probably depends on the preferential removal of such grains from soils. The particles must be transported only during the wet periods, that is during thaw or after heavy rains, since during the visits to the cave its walls were covered only by drops of condensation water and were devoid of seeping water. The episodic supply of material results in a visible lamination of microstromatolite (Fig. 5C). The microbial biofilm covering the vertical or even overhanging cave walls trap and bind the transported detrital particles (Figs 6C, D, 8D). The capacity to stabilize sand and finer grains has been experimentally confirmed (Meadows et al., 1994; Westall & Rincé, 1994; Dade et al., 1996). The particles are stabilized on the surface of microstromatolite and then covered with a newly developed biofilm (Fig. 6D). Simultaneously, an older part of the biofilm disintegrates due to senescence, which is probably facilitated by covering mineral particles. The laminae which comprise outsized quartz grains mark the especially wet episodes when relatively coarse-grained material could have been remobilized from soils and washed down into the cave (Fig. 5C). The microterracettes form in zones of more intense water flow (Fig. 3A–C). It is confirmed by their preferential formation along vertically oriented convex bends of the overhanging cave wall, thus the part where water flow is concentrated due to the surface tension. The downward inclined shape of particular microterracettes most probably results from plastic deformation and creeping of soft microstromatolite under the action of gravity (Fig. 3C).

The influx of seepage water is also important for the micro-organism assemblage forming the microstromatolites, which most probably depends on the input of organic matter from the surface (cf. Groth & Saiz-Jimenez, 1999). On the one hand, the limited energy of seeping water sorts

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**Fig. 8.** A – Three-dimensional porous network composed mainly of morphotype 6 (Actinomycetes); B – Detailed view of the network, spinose-walled morphotype 6 dominate, smooth-walled morphotype 7 occur subordinately (arrows), C – ovoid morphotype 5 (arrows) associated with spinose-walled filamentous morphotype 6; D – minute coccoid bodies of morphotype 4 (spores of Actinomycetes; small arrows) associated with filamentous morphotypes 6 and 7 (big arrow), in the centre platy mineral grain entrapped within organic filaments (EDS chemical composition is indicated); all photographs under SEM.
mineral grains and controls the fine-grained composition of microstromatolites. On the other hand, it allows delicate microbial biofilm to exist on cave walls; the higher energy of flow would cause deformation and scrubbing off of the microbial biofilm.

The lack of precipitation of minerals is most probably connected with the chemistry of the seeping water. Although direct data are lacking, we can hypothesize that the water in W Sopotni Wielkiej Cave is similar to the water in other non-karst caves in the flysch rocks of the Outer Carpathians. Zawierucha et al. (2005) reported that the vadose water in those caves is only slightly more mineralized than the rain water. The mean concentration of the Ca ion is only 9.5 mg/l, that is definitely lower than in karst caves. For example, the Ca content in water from selected Slovak karst caves ranges between 44.1 and 132.1 mg/l (Motyka et al., 2005) whereas in the vadose zone water of karst caves on the Kraków–Wieluń Upland it is from 52 to 137 mg/l (Różykowski, 2006). Moreover, one can suppose that the water which quickly percolates down after heavy rains, as it is in the discussed case, has lower concentration of ions due to its lower residence time (see discussion in Musgrove & Banner, 2004). Thus, the water in the studied cave most probably does not achieve the appropriate saturation to precipitate carbonate minerals.

The micro-organisms detected within the microstromatolites have been reported from other caves and have been supposed to influence – actively or passively – precipitation of minerals. Living bacteria belonging to Arthrobacter were detected within growing moonmilk deposits (Gradziński et al., 1997b) and cave pearls (Gradziński, 2000). Phosphatized microbial cells assigned also to this genus are reported from the speleothems of Grand Cayman Island (Jones, 2009b). Mineralized, commonly calcified, Actinomycetes, including various species of Streptomycetes, are known from cave deposits of Spain (Cañaveras et al., 2001), Italy (Groth et al., 2001), USA (Melim et al., 2008) and the Cayman Islands (Jones, 2009a, b). This leads to the suggestion that the mineralization of micro-organisms in the cavern environment is strongly dependent upon chemistry of the feeding water.

The described origin of the studied siliciclastic microstromatolites may shed some light on the formation of other cave deposits. The unconsolidated accumulations of fine-grained clastics on cave walls, including vermiculations common in many caves, can be also formed by the trapping and binding of detrital particles by micro-organisms. Several forms of speleothems recognized in some caves of the Polish Outer Carpathians may also share their origin with the discussed microstromatolites. They are also composed of siliciclastic material, display lamination and contain some laminae rich in coarser quartz sands (Urban et al., 2007).

The described stromatolites have a low potential for preservation. The only one possibility for their preservation is a quick cementation with calcium carbonates or opal acting as cement or covering a stromatolite as a younger flowstone (cf. Urban et al., 2007). Thus, clastic-rich layers occurring within flowstones and other speleothems (see Dzidzio et al., 1993; Gradziński et al., 1997a; Turgeon & Lundberg, 2001) may be fossil counterparts of microstromatolites formed during the break of crystallization and intense clastic supply. Since the micro-organisms entrapped detrital grains on the vertical and overhanging walls in the studied case, it seems plausible that they also play a role in stabilization of such grains deposited on a flowstone surface and later cemented with calcite or aragonite. Non-mineralized micro-organism cells have been decomposed and have not been preserved. Hence, the layers in question lack any traces of micro-organisms.

The closest modern counterparts of the described forms, both in morphology and origin, are sand ripples developing on steep sandstone crags described by Pentecost (1999) from Kent. They are composed of sand grains stabilized by algae and mosses and display relief strikingly similar to that of the described microstromatolites. The most important difference between the relief of sand ripples and the cave microstromatolites is a slightly greater lateral distance between neighbouring ripples (5.6–8.0 mm for the sand ripples). Other surface counterparts are litter dams occurring on slopes. They owe their origin due to accretion of small organic and mineral particles and their stabilization by mosses (Eddy et al., 1999). The spatial arrangements and relief of the litter dams are different, mainly because of their formation on the inclined, not vertical slopes.

Interestingly, a relief analogous to the described microstromatolites characterizes surfaces of several actively growing continental deposits. The surfaces of some cave flowstones and dripstones are crenulated (Hill & Forti, 1997, pp. 71, 105) or rippled (Ford, 1988). Similar ripples are formed on the growing icicles (e.g., Ogawa & Furuwaka, 2002). The cave draperies also display serrated edges composed of the stepped overhanging terracettes comparable to the terracettes formed by the described microstromatolites. Identical terracettes are composed of moonmilk deposits (Gradziński & Radomski, 1957). On the steeply inclined or overhanging walls their upper surfaces are downward inclined similarly to the terracettes of the described microstromatolites (Fig. 3E). Such orientation probably resulted from some instability of moonmilk having a pasty consistency and from its tendency to creeping down, which is in common with the discussed microstromatolites. Also travertines form similar regularly spaced terracettes. The lateral distance between the neighbouring travertine terracettes becomes shorter on steeper slopes (Hammer et al., 2010). Hence, the travertine terracettes developed on vertical and steep slopes may serve as a counterpart of the described ripples and terracettes formed by the microstromatolites. Nonetheless, in contrast with the microstromatolites and moonmilk examples, the rim of travertine terracettes is almost always perfectly horizontal (Fig. 3C; Hammer et al., 2010). It most probably results from the robust consistency of travertine deposits. Surprisingly, although all the above mentioned calcareous deposits originated mainly by the precipitation of crystals, not by the accumulation of detrital particles, they share their shape and geometric pattern with the discussed microstromatolites whose growth is governed by different factors. It adds a new dimension to the discussion on the factors influencing the shape and spatial arrangements of terracettes in recently growing travertines and speleothems.
CONCLUSIONS

1. The walls of W Sopotni Wielkiej Cave are covered with siliciclastic microstromatolites constructed by bacteria and Actinomycetes, which trap and bind mineral particles washed into the cave from overlying soil during wet episodes.

2. Newly precipitated mineral phases seem not to contribute to the microstromatolite growth.

3. The microstromatolites form ripples and microterraces on vertical and overhanging cave walls. Their shape depends upon the relief of a cave wall. The microterraces are located where water flow is more intense, mainly along vertically oriented convex bends of a cave wall.

4. It seems possible that trapping and binding mechanism influences the origin of clastic-rich layers within cave flowstones. Clastic grains were stabilized by micro-organisms and later were cemented with calcite or aragonite. The micro-organisms were subsequently decomposed and have not been preserved till now.

5. Sand ripples developing on almost vertical sandstone crags are a close genetic and morphological analogue of the cave siliciclastic microstromatolites.

6. Ripples (crenulations) on the speleothem and icicle surfaces, as well as stepped microterraces in speleothems and travertines share the same relief with the described microstromatolites in spite of their different origin.

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REFERENCES
