

Microbial mediation of stromatolite formation in karst-water creeks

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Abstract

Epilithic and endolithic biofilms were found to control the formation of stromatolites in karst-water creeks. We used microsensors to determine the influence of biological processes on chemical conditions within the microenvironment of crystal nucleation sites: the stromatolite surface. Phototrophic members of the biofilms consisted of mainly cyanobacteria and diatoms. Oxygen, pH, calcium, and carbonate concentration microprofiles at the stromatolite surface and boundary layer showed a strong diurnal rhythm of calcium carbonate precipitation. During illumination, photosynthesis caused oxygen production, a marked increase in pH and CO_3^{2-} concentrations, and a decrease in Ca^{2+} concentration at the stromatolite surface due to calcium carbonate precipitation. The opposite occurred in the dark, indicating decalcification. Calcite was approximately 16 times oversaturated in the bulk water, photosynthesis induced an increase of the supersaturation to >27 at the stromatolite surface under illumination, and respiration induced a decrease of the supersaturation to <10 in the dark. Photosynthetically stimulated calcium carbonate precipitation was confirmed by radioactive isotope ($^{45}\text{Ca}^{2+}$) uptake studies. Over a 24 h light : dark cycle, biofilms showed net calcification. Biotic activity within the stromatolite has a large effect on conditions at its surface and, therefore, contributes considerably to the stromatolite precipitation process.

Calcareous stromatolites are amongst the oldest known biological formations, and they provide insight into early Earth environments and climates. For this reason, it is essential to understand the processes governing their formation and dissolution. Presently forming stromatolites are rare in marine settings, but they are a common and much investigated feature of karst regions (Grüniger 1965; Merz-Preiss and Riding 1999). Such karst-water creek stromatolites have been termed tufa stromatolites, and they are defined as macroscopically laminated benthic microbial deposits produced by precipitation of minerals on organic tissue (Riding 1990). The role of biofilms in the formation of these stromatolites is still under debate. Early studies suggested that precipitation was primarily caused by CO_2 assimilation by cyanobacteria, eukaryotic algae, and plants (Pia 1926, 1933; Wallner 1934). Later hydrochemical investigations, however, concluded that precipitation is largely physicochemically driven by rapid CO_2 degassing from high- $p\text{CO}_2$ groundwater (Usdowski et al. 1979; Herman and Lorah 1987; Merz-Preiss and Riding 1999), with only a minor contribution from photosynthetic CO_2 removal. This conclusion was reached when it was observed

that calcium and dissolved inorganic carbon (DIC) were lost from water as it moved downstream, but no diurnal pattern was observed in creek-water chemistry. Even in instances where phototrophic communities were observed to affect whole-stream inorganic carbon dynamics (Spiro and Pentecost 1991), the development of tufa travertine deposits was seen as a largely abiotic process. The biofilms in the creeks investigated in the current study were very thin ($<100\ \mu\text{m}$), and indeed whole-stream-water chemistry parameters were hardly affected by biological activity. We tested the hypothesis that in these thin biofilms, strong shifts in local water chemistry are possible due to photosynthetic and respiratory activity, and that these shifts have highly localized effects on the stromatolite surface, where calcium carbonate precipitates.

Precipitation of calcite is initiated when calcium carbonate becomes supersaturated and suitable nucleation sites are present. Karst waters have a low $\text{Mg}^{2+} : \text{Ca}^{2+}$ ratio (<2), and, therefore, low-Mg-calcite is usually the main component in their stromatolite formations (Irion and Müller 1968; Arp et al. 2001). Photosynthesis and respiration can have a large effect on carbonate chemistry. Photosynthesis removes CO_2 and shifts the carbonate equilibrium toward carbonate, thereby increasing calcite saturation state (Ω). Respiration increases CO_2 and therefore has the opposite effect. Consequently, photosynthesis can lead to calcification, while respiration can lead to calcite dissolution when Ω decreases to values below one. In other aquatic settings, such as hypersaline lakes (Ludwig et al. 2005) and marine sediments (Werner et al. 2008) microsensor studies have demonstrated the potential for phototrophic communities to remove large amounts of CO_2 and enhance calcification. For stromatolite-forming biofilms in freshwater settings, this has not yet been investigated, and the view still prevails

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that the primary mechanism behind stromatolite precipitation is CO_2 degassing (Pentecost 1995).

It is also thought that, in addition to metabolic activity, biofilm components, particularly acid-organic compounds, in extracellular polymeric substances (EPS) that can strongly bind Ca^{2+} influence CaCO_3 precipitation. EPS is a common component of biofilms (Decho 1990; Nealson 1997; Neu and Lawrence 1999) and may contribute to precipitation in two ways: (1) suitably arranged acidic EPS groups create a template for precipitation and promote nucleation upon saturation with Ca^{2+} (organomineralization), and (2) the decomposition of EPS liberates HCO_3^- and Ca^{2+} and thereby increases calcite oversaturation (Arp et al. 2003; Trichet and Defarge 1995; Decho et al. 2005; Dupraz and Visscher 2005).

Previous studies in the Deinschwanger creek (*see* Methods) (Arp et al. 2001) were inconclusive as to the exact role of the biofilm in stromatolite formation, and diurnal rhythms had not been observed in this creek (Arp et al. 2003) due to resolution limitations of the bulk-water analysis methods used. Previous results have not excluded the possibility that the phototrophic biofilms diurnally control the local, *i.e.*, on the stromatolite surface, water chemistry. We therefore performed both *in situ* and *ex situ* experiments to investigate the role of the biofilm in stromatolite formation in more detail. We used fine-scale microsensor and β -imaging techniques to elucidate the influence of biofilm activity on carbonate chemistry and precipitation at the stromatolite surface.

Materials and methods

Creek sites—The Deinschwanger creek is located near Nürenberg, in southern Germany (49°23'N, 11°28'E), and the Westerhöfer creek is located near the Harz Mountains in northern Germany (51°45'N, 10°5'E). Both sites have been used for previous studies on calcification and are described in more detail in Arp et al. (2001) and Usdowski et al. (1979). Both sites show active laminated stromatolite formation of up to 1.8 mm yr^{-1} .

Sampling—Water samples for DIC and calcium measurements were taken from both the creeks and the aquariums during incubation. Fifteen-milliliter samples were filtered (0.2 μm) into exetainers (Labco Ltd.) containing HgCl_2 , sealed without air bubbles, and stored in the dark at 4°C until analysis.

Stromatolite samples for laboratory incubations were taken with a motorized core-drilling device in October 2005 and 2006 from Deinschwanger creek and May 2006 from Westerhöfer creek. Samples were stored in ambient creek water in coolers and transported to the laboratory within 24 h. At Deinschwanger creek, samples were collected from two distinct sites. Site 1 was a small stromatolite cascade downstream of the spring in a shaded site in the forest. The stromatolite was laminated, and it was several centimeters thick. At this site, the creek was $\sim 1\text{--}5$ cm deep and exhibited a fast flow rate of ~ 0.7 m s^{-1} . Site 2 was a well-illuminated, ~ 20 -cm-deep section of the lower creek, and it had a lower flow rate (~ 0.05 m s^{-1}) than site 1. Site 2

exhibited thin (up to 1 mm), fragile carbonate crusts. At the Westerhöfer creek, samples were taken from only one site, which was similar to site 1 at the Deinschwanger creek. Here, the creek was fast flowing and shallow, with laminated stromatolite several centimeters thick. Ambient pH at all sampling sites was 8.4.

In the laboratory, samples were stored in temperature-controlled (10°C), aerated recirculating aquariums (total volume ~ 30 liters) containing creek water. Creek water was collected without air bubbles, in 20-liter plastic containers, at the same time cores were taken. The water was stored in the dark at 4°C until it was added to the aquarium reservoir. Samples were illuminated at 150 mol quanta $\text{m}^{-2} \text{s}^{-1}$ on a 12 h light : dark (LD) cycle.

Water analyses—Bulk water pH was determined with a MA130 pH electrode (Mettler Toledo) directly in the creek and in the experimental aquaria. Ca^{2+} was determined by inductively coupled plasma–optical emission spectrometry (ICP-OES; Perkin Elmer Optima 3300 DV), and DIC was determined by CO_2 coulometer (UIC Coulometrics).

Algal pigment analysis—Triplicate biofilm samples from Deinschwanger creek (sites 1 and 2) and single samples from incubation experiments were analyzed for algal pigments to assess the change in algal communities induced by experimental conditions. Samples taken freshly from the creek were stored on ice and frozen (-20°C) as soon as possible (<24 h). Samples used for laboratory studies were immediately frozen after the incubations. For analysis, frozen samples were placed in 15-mL centrifuge tubes, in 3 mL of 100% acetone, and sonicated on ice for 10×1 min. Pigments were extracted from the mixture at -20°C overnight. Samples were then filtered (Acrodisc CR syringe filter, 0.45- μm pore size, PALL GmbH, Dreieich, Germany, Gellman Laboratory) and diluted with water to a final concentration of 70% acetone. Photopigments were separated on a high performance liquid chromatography (HPLC) (2690 separation module, Waters), equipped with a Eurospher-100 C18, 5- μm Vertex column (Knauer), according to the method of Wright et al. (1991). Absorption spectra of the separated compounds were measured on a Waters 996 photo-diode-array detector, and pigments were quantified and identified by comparison to pigment standards (DHI Waters and Environment).

Microelectrode measurements and calculations—Microelectrodes are ideal tools to measure chemical gradients in biotic systems at fine (μm) scales (*see* Gieseke and de Beer [2004] for a review of microsensor form, function, and application to microbiological samples). Microelectrode profiles of pH, O_2 , and Ca^{2+} concentration gradients were measured *in situ* at the two creeks. Profiles of these and CO_3^{2-} were also made *ex situ*, in the laboratory, using biofilm-covered stromatolite cores.

Liquid-membrane Ca^{2+} and pH sensors (tip = 20 μm) were prepared and calibrated as described previously (de Beer et al. 1997, 2000). Liquid-membrane CO_3^{2-} sensors were constructed similarly to the other liquid-membrane sensors and calibrated as described in de Beer et al.

(unpubl.). For carbonate system components, pK1 (6.464) and pK2 (10.489) were calculated after Millero (1979), for freshwater and at 10°C. Fast-responding O₂ microsensors (tip = 10 μm) were prepared as described previously (Revsbech 1989). Biofilms were very thin (50–100 μm), and it was difficult to distinguish the biofilm surface from that of the calcium carbonate tufa. Indeed the biofilms were partly endolithic. The glass electrodes could not penetrate the hard mineral surface. All electrodes were, therefore, placed at the stromatolite–biofilm surface while the sample was viewed through a dissection microscope. The surface was then set at 0, and all measurements were made above the surface. Negative distances indicate that the sensor was above the stromatolite–water interface, that is, in the overlying water. Sensors were connected to a micromanipulator that was fixed onto a motorized stage (VT-150, Micos), which allowed reproducible positioning of the sensor tip with 1 μm precision. The microelectrodes were connected to a picoammeter (O₂ electrode) or a millivoltmeter, and the meter output was collected by a data-acquisition device (NI-Daq 6015, National Instruments). After positioning at the surface, profiling was done automatically; motor control and data acquisition were performed with a computer and custom-written software (*μ*-Profiler, Dr. L. Polerecky). All profiles were corrected for offset to ion levels in the bulk liquid, as measured by the water analyses. For in situ microelectrode measurements, the micromanipulator was mounted to a heavy stand and placed in the creek, on the downstream side of the measuring position.

Interfacial fluxes (*J*) were calculated from the concentration profiles using Fick's first law:

$$J = D \times (dc/dx) \quad (1)$$

where *D* is the diffusion coefficient, and *dc/dx* is the interfacial concentration gradient, i.e., the concentration gradient in the mass boundary layer directly adjacent to the stromatolite surface. The diffusion coefficients of O₂ and Ca²⁺ are literature-derived values corrected for temperature and type of counterion (Broeker and Peng 1974; Li and Gregory 1974). Estimated diffusion coefficients of oxygen are, 4°C: $1.28 \times 10^{-9} \text{ m}^2 \text{ s}^{-1}$; 10°C: $1.57 \times 10^{-9} \text{ m}^2 \text{ s}^{-1}$; and 17°C: $1.934 \times 10^{-9} \text{ m}^2 \text{ s}^{-1}$. Diffusion coefficients for Ca²⁺ with HCO₃²⁻ as counterion are, 4°C: $0.546 \times 10^{-9} \text{ m}^2 \text{ s}^{-1}$; 10°C: $0.67 \times 10^{-9} \text{ m}^2 \text{ s}^{-1}$; and 17°C: $0.827 \times 10^{-9} \text{ m}^2 \text{ s}^{-1}$.

Calcite saturation is given by the saturation-state omega,

$$\Omega = [\text{Ca}^{2+}] \times [\text{CO}_3^{2-}] / K_{(\text{calcite})} \quad (2)$$

where the numerator is the ion concentration product (ion activities were not incorporated), and the denominator the solubility product. $\Omega > 1$ indicates supersaturation, while $\Omega < 1$ indicates undersaturation. The solubility product for calcite was taken from Plummer and Busenberg (1982).

Incubation with radiotracers—Stromatolite samples from both creeks were subject to radiotracer incubation with ⁴⁵Ca²⁺. Stromatolite samples were incubated under three experimental conditions: (1) under illumination at 150 mol

quanta m⁻² s⁻¹, (2) in the dark, and (3) fixed with formaldehyde (2%) control. Samples were analyzed by both scintillation counting and β-imaging. During each treatment, stromatolite samples were incubated in 400 mL of filtered creek water at 20°C and gently stirred. Prior to addition of the radiotracer, the stromatolite pieces were left for at least 2 h to recover from the transfer. The radioactive tracer, ⁴⁵Ca²⁺ as CaCl₂ (Amersham Pharmacia Biotech), was added to each beaker to a final activity of 3 kBq mL⁻¹ (specific activities were 74.0 MBq mL⁻¹) in all the experiments. Samples were incubated with tracer for 1, 3, 6, and 24 h.

After the incubation, the samples were fixed in 2% formaldehyde for 2 h. The fixed stromatolite samples were then washed four times for 5 min in 200 mL of filtered creek water. Samples were then left to air dry overnight.

Samples for scintillation counting were weighed and dissolved in 10 mL of 12 N HCl. Two milliliters of dissolved sample were then added to 3 mL of Ultima gold scintillation cocktail (Perkin-Elmer), and radioactivity was determined using a Packard TR 2500 liquid scintillation counter operating in efficiency-tracing mode to correct for quench.

Samples for β-imaging were immersed in Epothin resin (Buehler), cut into ~4-mm cross sections with a diamond band saw (Proxon MBS 240E), and polished flat with a Buehler Beta polishing system. β-imaging autoradiographs of the stromatolite samples were acquired in a Micro-Imager (Laniece et al. 1998). The resin fixation and polishing process occasionally produced artifacts in the data obtained by the β-imager. These artifacts were caused by air bubbles in the resin, which filled with radioactive dust when the samples were cut and polished. They were easily recognizable as a circular “hotspot” that coincided with an air-bubble-induced hole. These artifacts were removed from the analysis with Photoshop CS (Adobe Systems Inc.).

Results

Sample description and pigment analysis—Stromatolite biofilms in both the Deinschwanger and Westerhöfer creeks were green-brown and composed of cyanobacteria, diatoms, and associated nonphototrophic bacteria. Pigment composition was analyzed only for samples from Deinschwanger creek, site 1 (small tributary on northern valley side) and site 2 (lower end section, near end of stromatolite deposition).

Samples from site 1 consisted of thick stromatolite cores, the tops of which were covered by an ~50–100-μm-thick green and brown biofilm, while those at site 2 consisted of an equally thick, brown-colored biofilm covering a 1-mm-thick, fragile carbonate crust. The appearance of the samples did not change markedly during the study, but biofilm thickness sometimes increased slightly. Pigment composition did not change markedly during incubation (Fig. 1). The pigment concentrations and compositions in samples at the end of incubations were similar to those from the creek, showing that the algal community remained stable during incubations. Pigment concentrations were higher at site 2. The compositions of the pigments were similar at both sites. Pigments were dominated by chlorophyll *a* and β-carotene. Smaller amounts of fucoxanthene, zeaxanthine, phaeophy-

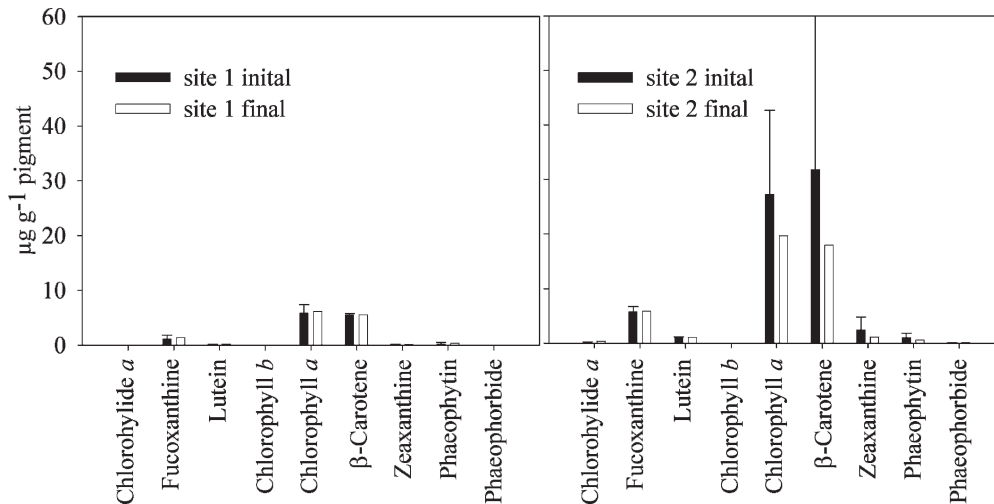


Fig. 1. Algal pigments in tufa samples from Deinschwanger creek (sites 1 and 2) prior to and following incubation (initial samples $n = 3$; final samples $n = 1$).

tin, and chlorohyllide α were also seen in both sites. Fucoxanthene, zeaxanthine, and phaeophytin indicate the presence of cyanobacteria and diatoms in these biofilms.

In situ measurements—In situ profiling was complicated by the heterogeneous substrate topography, debris flowing down the river, and difficult visualization of the hard substratum under poor light conditions. Consequently, in situ microprofiling was often interrupted by breakage of microsensors. In situ concentration-gradient profiling was successfully carried out for O_2 , pH, and Ca^{2+} in both Deinschwanger and Westerhöfer creeks, although not all

species could be profiled both in the day and night at both sites (Figs. 2, 3). The data clearly indicate a day to night change in chemical conditions on the stromatolite surface. Fluxes calculated from profiles are given in Table 1. Photosynthesis occurred at the stromatolite surface of both creeks in daylight and caused a flux of O_2 from the biofilm to the water column, an increase in pH at the stromatolite surface, and a concomitant decrease in Ca^{2+} ion concentration. The night profiles showed the opposite. In Westerhöfer creek, a decrease in O_2 concentration occurred with respiration, and an increase in Ca^{2+} concentration was evident. At Deinschwanger creek, the only pH profile was obtained at night, and it showed a decrease in pH at the stromatolite surface, indicating respiratory activity. The efflux of Ca^{2+} in the dark was lower than its influx in the light; thus, during a 12 : 12 h photoperiod, there was a net

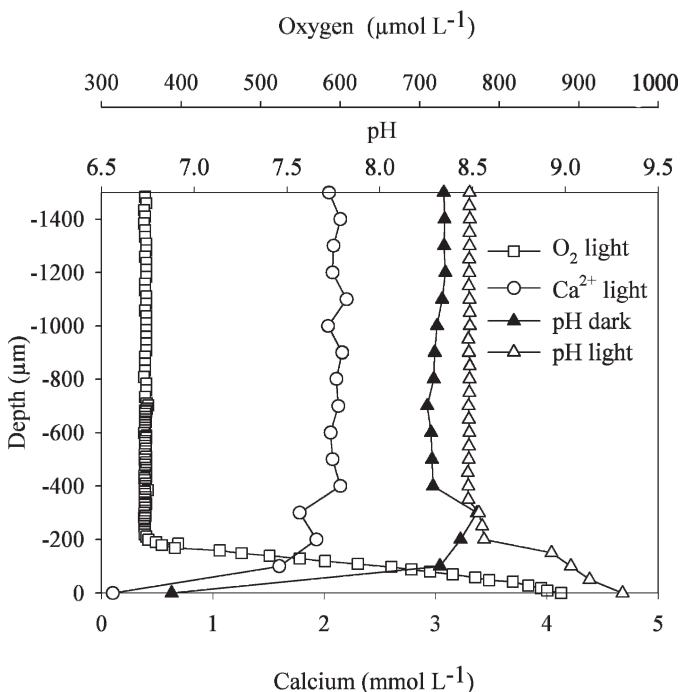


Fig. 2. In situ concentration profiles of Ca^{2+} , pH, and O_2 at Deinschwanger creek.

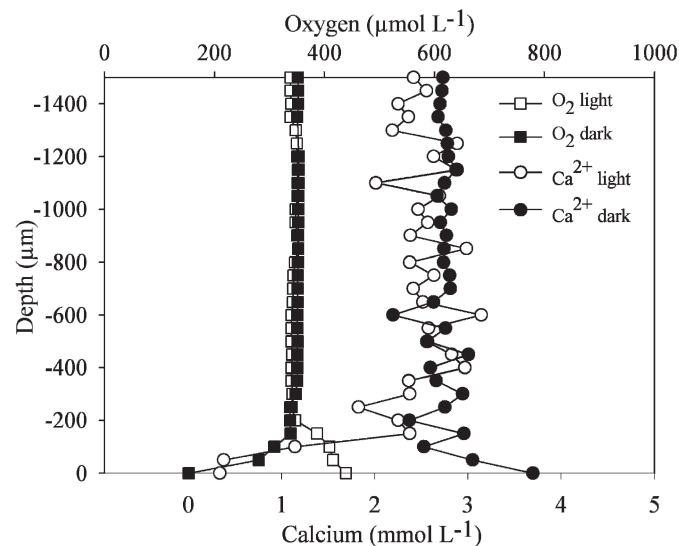


Fig. 3. In situ concentration profiles of Ca^{2+} and O_2 at Westerhöfer creek.

Table 1. In situ fluxes ($\text{mol m}^{-2} \text{s}^{-1}$) of oxygen and calcium in Deinschwanger and Westerhöfer creeks. A negative flux indicates a flux from the water column toward the tufa surface.

	Deinschwanger creek		Westerhöfer creek	
	Oxygen	Calcium	Oxygen	Calcium
Day	5.6×10^{-6}	-2.0×10^{-5}	1.0×10^{-6}	-1.3×10^{-5}
Night			-9.1×10^{-7}	7.3×10^{-6}

flux of $1.2 \times 10^{-5} \text{ mol Ca}^{2+} \text{ m}^{-2} \text{ s}^{-1}$ toward the stromatolite surface over a full 24-h cycle.

Ex situ measurements—The in situ measurements were complemented with a detailed ex situ study. In samples from both creeks, under illumination, O_2 and pH increased due to photosynthesis, coincident with a decrease in Ca^{2+} . The opposite occurred under dark conditions (Figs. 4, 5). Respiration resulted in O_2 consumption and a decrease in pH. The Ca^{2+} profile also changed with this change in metabolism, such that Ca^{2+} uptake (CaCO_3 precipitation) ceased, and its release from the tufa (apparent CaCO_3 dissolution) occurred. The carbonate concentration increased at the stromatolite surface during light conditions, and carbonate concentration decreased in the dark (Fig. 5). Fluxes calculated from the profiles are shown in Table 2. The fluxes of both O_2 and Ca^{2+} were similar during in situ measurements and the laboratory incubations. Accurate quantitative comparisons were difficult because of the heterogeneity of the samples.

Carbonate profiles, measured by carbonate microsensors in several samples from Westerhöfer creek, agreed well with carbonate profiles modeled from the bulk-water DIC concentrations and pH microprofiles using the CO_2 system equations (Zeebe and Wolf-Gladrow 2001) (data not shown). Subsequently, bulk-water DIC and pH microprofiles were used to predict the CO_3^{2-} concentration and Ω profiles for tufa samples (Fig. 6). These calculations revealed an increase in CO_3^{2-} in light, which caused Ω to

become >20 . The opposite occurred in the dark, when CO_3^{2-} and Ω became <10 .

Water flow had a small effect on the photosynthesis rates. Increasing the flow rate from 0.009 L s^{-1} to 0.035 L s^{-1} caused the diffusive boundary layer (DBL) to decrease in thickness from $\sim 600 \mu\text{m}$ to $200 \mu\text{m}$ and the areal rate of O_2 production to increase from 1.93×10^{-6} to $2.47 \times 10^{-6} \text{ mol m}^{-2} \text{ s}^{-1}$. The height of the DBL indicates the range over which diffusion determines chemical gradients. The smaller the DBL, the faster chemicals can move between the overlying water and the sediment surface. Flow rates in situ were very fast, and the DBL was probably slightly less than that in our ex situ experiments. This, however, does not change the interpretation of the clear diurnal calcium dynamics observed.

Radiotracer incubations—Scintillation counts of whole samples showed that calcium was incorporated under all conditions, but rates of incorporation in the light were approximately twice as high as in the dark and controls (Fig. 7). Dark rates equaled control rates, indicating that in the dark, the calcium uptake was mainly isotopic exchange. The β -imaging (Fig. 8) confirmed that uptake of $^{45}\text{Ca}^{2+}$ was stimulated by light, and that Ca uptake preferentially occurred at the stromatolite surface. Indeed, $^{45}\text{Ca}^{2+}$ uptake appears to have occurred most strongly in the middle of the biofilm, but it could be seen throughout the whole biofilm (photosynthetic zone), not only at the biofilm-

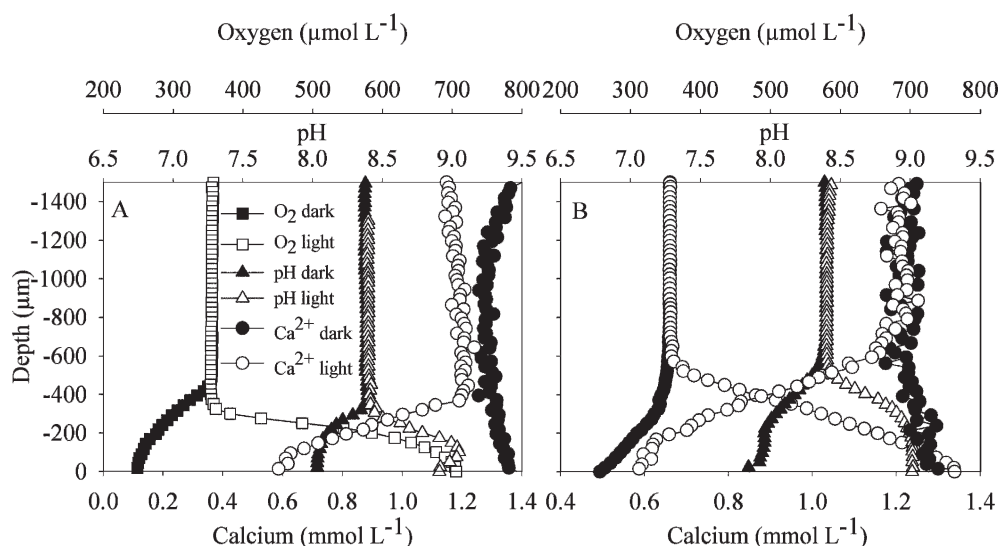


Fig. 4. Ex situ concentration profiles of Ca^{2+} , pH, and O_2 from Deinschwanger creek tufa biofilm samples: (A) site 1, (B) site 2.

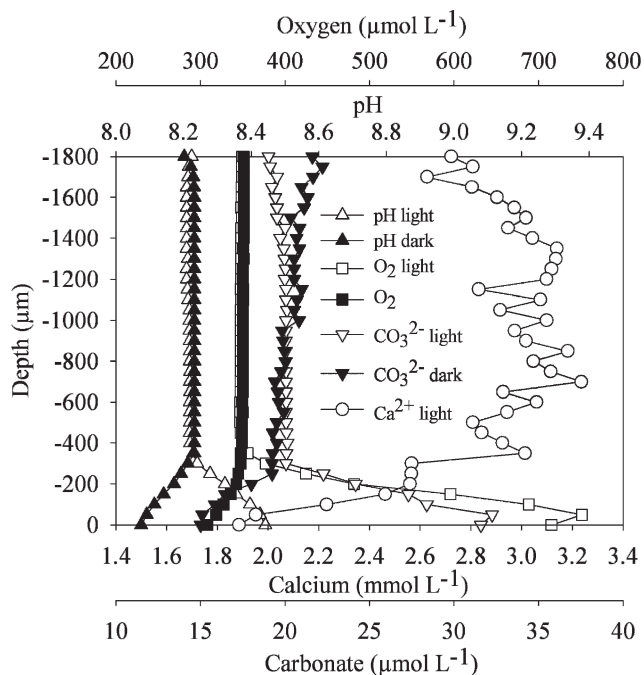


Fig. 5. Ex situ concentration profiles of Ca^{2+} , pH, O_2 , and CO_3^{2-} from Westerhöfer creek tufa biofilm samples.

overlying water interface. In the dark and control samples, radioactive signals were detected throughout the whole sample.

Discussion

There has been a long controversy regarding whether stromatolites in karst-water creeks are formed by biological removal of CO_2 , so by photosynthesis, or by CO_2 degassing. It is, however, conceivable that both processes occur simultaneously in the creek: in the water column, the carbonate system is driven by abiotic forces, as suggested previously, while stromatolite formation is driven by metabolic processes. We propose that the most dominant calcification process in the creek does not result in stromatolite formation, but it occurs in the water column, and the precipitated calcium carbonate is washed from the spring area as suspended crystals. This calcification process is driven by CO_2 degassing and manifests itself as a downstream calcium loss from the water column, and a downstream pH increase, with no diurnal pattern to the calcium loss, stream pH, or saturation-state parameters.

This physical process has been well documented in similar systems, and it was also found in the creeks of this study (Usdowski et al. 1979; Merz-Preiss and Riding 1999).

Parallel to this physicochemical process, biologically controlled calcification results in stromatolite formation. Local CO_2 fixation by photosynthesis leads to calcification in the top 100 μm of the stromatolite surface. Interestingly, apparent decalcification is observed during the night, due to respiration; so, locally, the biological processes can counter the massive physical process of outgassing. The biological processes exert no significant influence over the whole stream carbonate system, and so they do not induce diurnal rhythms in pH and Ca^{2+} concentrations in the water column. Indeed, a rough, conservative calculation can be made regarding the relative contributions of the two processes. Assuming about 100 m^2 of stromatolite area, a flow rate of 2 L s^{-1} , Ca^{2+} flux estimated from actual tufa deposits ($\sim 4\text{-mm}$ -thick annual lamination, unpubl. data), and Ca^{2+} loss from the bulk water of 0.4 mmol L^{-1} , stromatolite precipitation accounts for less than 15% of the creek's Ca loss. The biological processes are only measurable at the stromatolite surface with high-resolution methods, such as microsensors. The biological effects on the water-column pH or Ca^{2+} concentrations are lost in the overwhelming background of the larger CO_2 degassing-driven process. However, locally, the effects of biological processes are highly significant. The biological precipitation process, which results in stromatolite growth, manifests itself as precipitation at the stromatolite surface and exhibits a clear diurnal rhythm, synchronous with biofilm photosynthetic and respiration processes. The present microsensor pH, O_2 , and Ca^{2+} profiles clearly demonstrate this diurnal pattern and, therefore, a decisive biological control on the microenvironment and Ca^{2+} removal for stromatolite formation.

It has also been argued that biofilms merely act as nucleation site for calcification or concentrate calcium by binding to EPS (Dupraz and Visscher 2005). With this in mind, two possible scenarios will be discussed in view of the observed Ca^{2+} microprofiles and radiotracer data: (1) Ca^{2+} uptake in newly produced EPS without precipitation and later Ca^{2+} release due to pH decrease or EPS decomposition causes precipitation in a second step, or (2) CaCO_3 precipitation is caused by photosynthetic CO_2 removal.

The first scenario does not explain our radiotracer data: EPS is produced mainly under illumination and decomposes in the dark (Decho and Lopez 1993; Epstein 1997; Staats et al. 2000). If release of Ca^{2+} from EPS were driving supersaturation under these conditions, CaCO_3 would be

Table 2. Ex situ fluxes ($\text{mol m}^2 \text{s}^{-1}$) of oxygen, calcium, and carbonate in Deinschwanger and Westerhöfer creeks. Numbers in parentheses identify site 1 or 2. A negative flux indicates a flux toward the tufa surface.

	Deinschwanger creek		Westerhöfer creek		
	Oxygen	Calcium	Oxygen	Calcium	Carbonate
Light	4.37×10^{-6} (1)	-1.30×10^{-6} (1)	3.24×10^{-6}	-3.42×10^{-6}	2.39×10^{-7}
	1.62×10^{-6} (2)	-8.88×10^{-7} (2)			
Dark	-3.75×10^{-7} (1)	1.16×10^{-7} (1)	-2.47×10^{-7}		-1.44×10^{-7}
	-4.3×10^{-7} (2)	0 (2)			

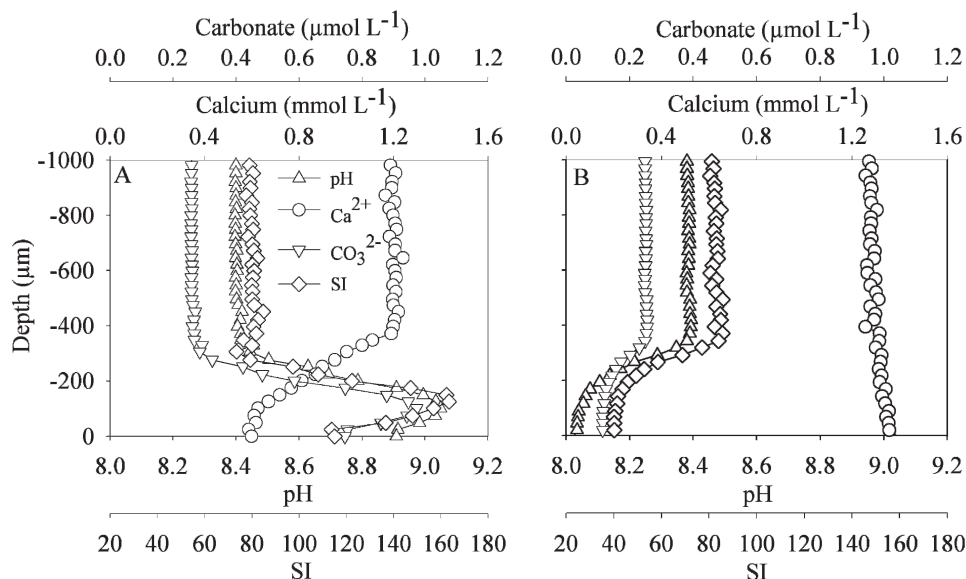


Fig. 6. Measured microprofiles of pH and Ca^{2+} in the (A) light and (B) dark and predicted CO_3^{2-} and calcite saturation Ω calculated from the measured pH profiles and bulk-water DIC concentrations. Samples are from Deinschwanger creek, site 1.

precipitated in the dark. All samples were illuminated immediately before the tracer experiment. If release of Ca^{2+} bound to EPS were initiating precipitation, $^{45}\text{Ca}^{2+}$ uptake should have been seen in both the light and dark treatments—in the light, as the tracer was bound to EPS, and in the dark, as previously bound Ca^{2+} was released, leading to CaCO_3 supersaturation and precipitation. This was not the case; $^{45}\text{Ca}^{2+}$ uptake was observed only in the light. We do not exclude that pH changes induced by photosynthesis–respiration cycles make EPS more amenable to Ca^{2+} binding in the light and release in the dark (Gregor et al. 1996). Such a scenario would show Ca^{2+} uptake during the day, due to binding to EPS, and release in the night, as EPS groups become unfavorable to binding. Under such a case of binding and release, the Ca^{2+} is not precipitated but merely removed from and returned to the

water phase. Since EPS can have only a limited number of binding sites, the uptake and release would be transient, occurring during the pH transitions upon illumination or darkening. No such behavior in Ca^{2+} microprofiles was observed: typical light and dark profiles were stable for 12 h. The Ca^{2+} taken up during the day was also clearly precipitated as CaCO_3 .

Thus, the second scenario is the most probable: biofilm photosynthetic activity changes conditions at the stromatolite–water interface (the site of precipitation), by removal of CO_2 , to such an extent that rapid CaCO_3 precipitation occurs.

As reported in Arp et al. (2001), the stromatolite biofilms were dominated by cyanobacteria and diatoms. The pigment data show that the community is stable when removed from the creek to the laboratory, and it is thus an

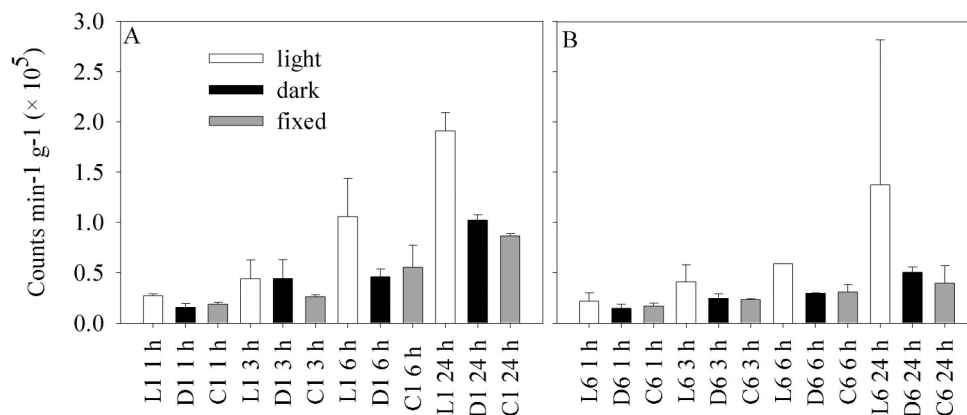


Fig. 7. $^{45}\text{Ca}^{2+}$ uptake of tufa samples from Deinschwanger creek: (A) site 1, (B) site 2. Open bars represent light incubations, black bars dark represent incubations, and gray bars represent formaldehyde fixed controls. X-axis labels refer to light (L), dark (D) and control (C) incubations as well as incubations time (h) (mean \pm SE, $n = 2$).

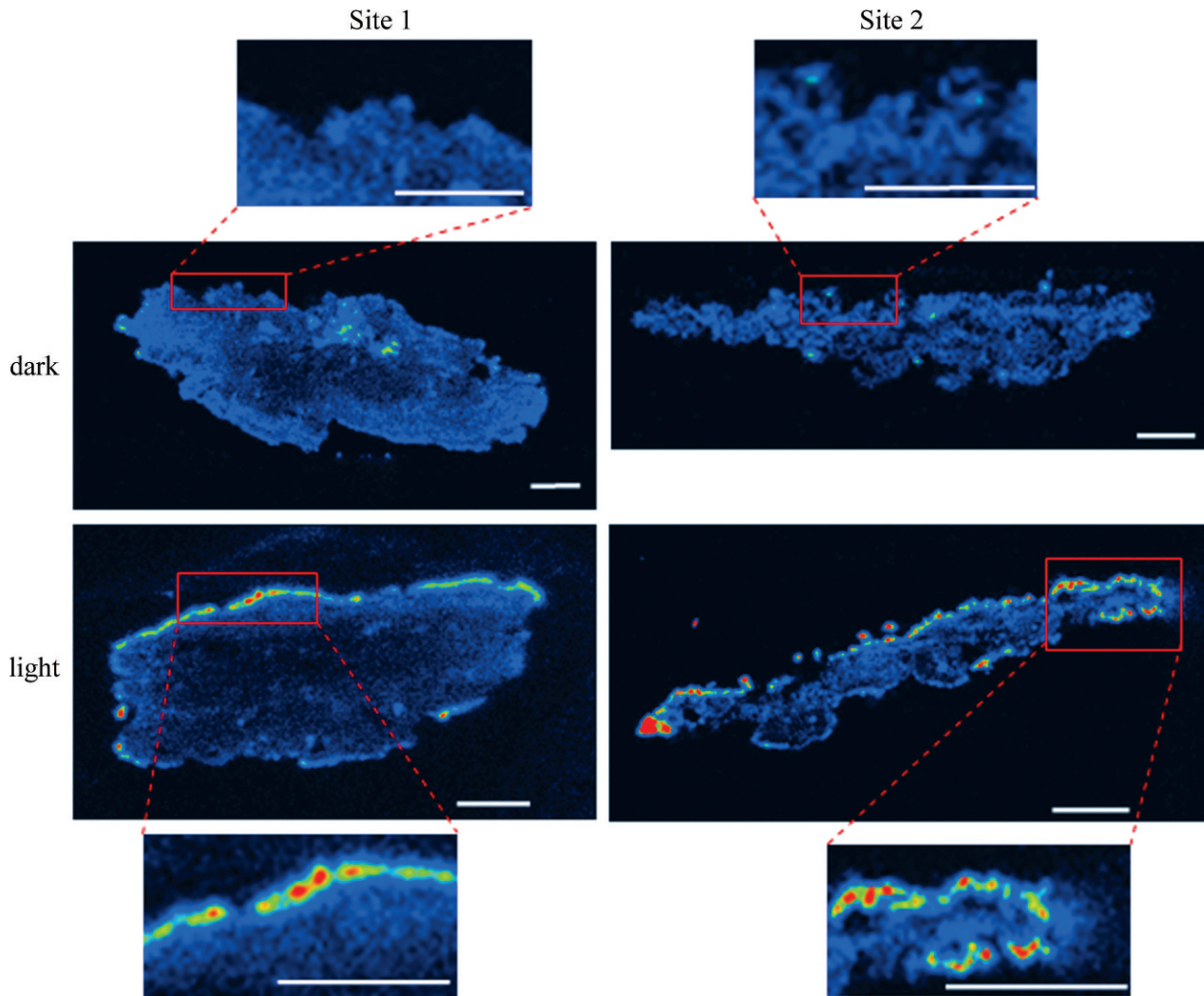


Fig. 8. β -images of light- and dark-incubated tufa samples from Deinschwanger creek after 24 h. The original biofilm surface is at the top of each sample. Scale bar = 3 mm.

appropriate model for ex situ experimental investigations. Water-chemistry parameters also varied somewhat between our in situ and ex situ measurements. The Ca^{2+} and DIC concentrations decreased as the incubations continued. Particulate deposits were observed on the base of the recirculating incubation system, but a light–dark pattern to calcium loss was only observed at the stromatolite surface. The loss of Ca^{2+} and DIC may be seen in the relatively low bulk-water Ω values observed during the light in our ex situ measurements (Fig. 6). Despite these variations, and although the resultant rates may differ somewhat between the in situ and ex situ measurements, general patterns remained the same. The differences in rates (which seem generally higher for in situ measurements) might be an effect of the different analytical setups (laboratory vs. creek), or they may be due to the natural variation caused by mat patchiness. The level of replication used for the in situ measurements was necessarily small; thus, quantitative extrapolation of the microsensor observations to whole-creek systems must be done with caution. The general agreement among the processes indicated and patterns

observed among the different measurements, however, makes it appropriate to discuss, in general terms, the overall processes occurring in the mats, based on both our in situ and ex situ observations.

The apparent, slight, Ca^{2+} flux to the water phase in the dark period, despite an Ω value always greater than 1, may be explained by dissolution that occurred below the stromatolite surface, in the endolithic part of the biofilm, which the microsensors could not reach. Heterotrophic activity probably continues below this zone and creates an environment conducive to stromatolite dissolution. As the stromatolite dissolves in the dark, Ca^{2+} diffuses upward, past the stromatolite surface, and is recorded by microsensors as Ca^{2+} flux from the stromatolite.

The importance of photosynthesis, not merely the presence of the biofilm and its potential nucleation sites, was also shown by the incubation of samples with the radioactive tracer $^{45}\text{Ca}^{2+}$. Incorporation rates of the tracer were highest in the photosynthesizing sample, and β -imaging showed that the tracer was specifically incorporated in the biofilm-covered surfaces. The incorporation

can be seen at or just below the surface of the light-incubated samples (Fig. 8). The region just below the surface is within the biofilm and would be the zone of highest pH. The incorporation seen occasionally at the base of the samples was not an artifact. The samples were incubated in aquariums for several weeks, and the biofilms had colonized parts of the underneath and sides of the samples. Thus, both the light-stimulated uptake rates and the distribution of tracer incorporation are compatible with photosynthetically driven calcification.

It is interesting that not only was the calcium flux toward the stromatolite surface higher in the *in situ* measurements, but so was the rate of photosynthesis. This agrees with laboratory results on flow rates, which suggest that photosynthesis should be lower in the slower flow rates of the laboratory.

Because replication in this study was low, the exact rates determined herein need to be taken with some caution, but the diurnal pattern and net precipitation (rather than net dissolution) over the light : dark cycle observed here were remarkably persistent over all samples.

While further studies are needed to elucidate the exact mechanism and rates of nucleation and precipitation in these stromatolite creek systems, the biological activity of their phototrophic communities appears to be much more important than previously thought. Despite not driving the total-stream carbonate-system dynamics, the biofilm has a strong influence over the state of the carbonate system at the sites of nucleation. We have demonstrated this, for the first time, by observing a strong diurnal pattern in Ca^{2+} and CO_3^{2-} concentration profiles and Ω , a pattern that coincides with the photosynthetic activity of the biofilm.

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