

Calcite production by the calcifying green alga *Phacotus lenticularis*

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ABSTRACT

The importance of carbonate precipitation by phytoplankton in fresh water lakes has not been sufficiently considered in global carbon cycles and climate change scenarios. The objective of this study was to determine the influence of the calcifying bivalved phytoflagellate *Phacotus lenticularis* (Ehrenberg) Deising 1866 on the total calcite precipitation in five European hard-water lakes. For this purpose, an accurate mass determination of single *Phacotus lenticularis* shells was required. We developed a novel methodological approach to precisely determine the volume and mass of the calcified shells. Focused ion beam (FIB) techniques were employed to investigate internal structural features. Thin layer cross-sections of the shell profiles were reproduced and perforation as well as the crystalline structure of the calcite plates were monitored. 3D-shell models were computed by 360° rotation of the shell cross-sections using a CAD 3D imaging software to calculate precise volumes and estimate realistic masses. In contrast to previous estimates, we determined a 2.8-fold higher shell mass of 0.86 ng CaCO₃ (SD=0.18) for the highly massive shells at a mean volume per individual of 334.1 μm³ (SD=70). An initial shell porosity of less than 5% was derived from thin layer cross-section images, resulting in a presumed mean shell density of 0.0026 ng μm⁻³. The shell diameter was significantly influenced by the lake's origin. The shells from each lake displayed substantial variations in diameter and shape. The pores in the shells showed two variations. Wider pore canals penetrated the whole shell wall, whereas small, elongated pores were located along the interspaces between calcite crystals with tabular habit. The approximate average dimensions of these calcite plates were 1.0×1.6×0.2 μm. The mean lateral wall thickness at the rim and centre of the shell were 1.98 μm (SD=0.42) and 0.79 μm (SD=0.17), respectively. The average carbonate precipitation by *Phacotus lenticularis* in relation to the total epilimnetic suspended calcite precipitation was 6% in the oligotrophic lake Großer Ostersee (Bavaria, Germany). During the growing season, *Phacotus lenticularis* contributed up to 21% of the particulate calcium carbonate in the epilimnion. These findings suggest that *Phacotus lenticularis* should be considered in the assessment of hard-water lake carbon cycling.

Key words: Autochthonous calcite precipitation; calcite shells, hard-water lake; mass determination; focused ion beam.

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INTRODUCTION

The role of inland aquatic systems in C-cycling is poorly understood. In addition to the terrestrial C-input into aquatic systems (Cole *et al.*, 2007), there can also be substantial C-fixation by primary producers within aquatic systems. Carbonate precipitation induced by planktonic algae is a process that has been widely investigated in marine environments. However, it also occurs in freshwater ecosystems, but there is less information available about its importance in these ecosystems. The calcifying freshwater flagellate *Phacotus lenticularis* (Ehrenberg) Deising 1866 (Chlorophyceae, Chlamydomonadales, Phacotaceae) is abundant in temperate hard-water lakes worldwide. These unicellular green algae have been reported to incorporate remarkable amounts of CaCO₃ in their shells (Giering *et al.*, 1990; Koschel *et al.*, 1987; Koschel and Raidt, 1988; Pocratsky,

1982; Sieminska, 1952; Sladeczek *et al.*, 1958; Steinberg and Klee, 1983), consisting of 98% - 99% pure CaCO₃ (Schlegel *et al.*, 1998). It is assumed that it temporally contributes up to 100% of the autochthonous calcite precipitation during mass development (Koschel and Raidt, 1988). Consequently this species brings about high autochthonous CaCO₃ inputs in these lakes and plays a definite role in limnic carbonate sedimentation as a sediment contributor (Koschel *et al.*, 1987; Müller and Oti, 1981).

Previous works have described the external and structural features of *Phacotus* shells (often named *lorica*) in great detail. The mineralised external housing has a simple radially symmetric morphology and is formed by rhombohedral calcite crystals that are arranged in several rings around a central rotational axis (Kamptner, 1950; Koschel and Raidt, 1988). However, several important aspects remain controversial. For instance, it is unknown

if the interspaces between the rhombohedral crystals are filled with calcite or remain empty. Also, internal orientation and extensions of the crystals have not been described so far. Moreover, there have been no explicit measurements of their porosity. *Phacotus* shells are likely to vary in morphotype and surface porosity over time and location (Schlegel *et al.*, 2000). Variations in shell diameter between 6-17 μm have been reported (Giering *et al.*, 1990; Koschel and Raidt, 1988; Pocratsky, 1982; Schlegel *et al.*, 2000; Steinberg and Klee, 1983). However, none of these works systematically determined the precise amount of calcite within a shell. Furthermore, there has been no explicit assessment of the individual variations of the general dimensions, nor calculations of the volume of *Phacotus* shells from different lakes.

A single provisional estimation based on conductometry has been reported by Koschel *et al.* (1987), which suggested a shell mass of 0.3 ng CaCO_3 per individual shell. This result was derived from a particularly suitable *Phacotus* mass development in which the shells accounted for 100% of autochthonous calcite. To the best of our knowledge, this value has not yet been verified, and a repetition of the measurement seems difficult because i) it is exceptionally rare that mass developments account for 100% of autochthonous calcite precipitation, and ii) in all other cases mass determination of exclusively *Phacotus* shells is generally impeded by co-precipitated calcite crystals.

In our study, we built the basis for a quantitative determination of the potential role of *Phacotus lenticularis* in carbonate precipitation in lakes. Our methodical approach revealed new insights on the internal ultrastructure of the *Phacotus* shell and enabled us to verify and estimate the porosity of the material. We measured the relevant external shell parameters, calculated the precise shell volumes, and estimated the realistic shell masses of 24 individual shells from five different lakes. We investigated if there is a variation of the *Phacotus* diameter

in lakes with different trophic states and during the growing season. Finally, we determined the amount of *Phacotus* calcite in relation to the total calcite precipitation in the epilimnetic layer of the oligotrophic lake Großer Ostersee (Bavaria, Germany) and addressed the question of whether *Phacotus lenticularis* significantly affects the CaCO_3 budget of this lake.

METHODS

Study sites

Phacotus shells for mass determination were derived from five lakes in southern Bavaria, Germany: lake Abtsdorfer See (ABS), lake Altmühlsee (ALT), lake Großer Ostersee (GOS), lake Hopfensee (HOP) and lake Igelsbachsee (IGS). The lakes differed in surface area, maximum depth, and trophic state (Tab. 1), but held significant *Phacotus lenticularis* occurrences with cell densities of more than 1×10^5 individuals per litre (ind. L^{-1}). Intensive investigations on suspended carbonate and *Phacotus* population dynamics in lake Großer Ostersee were conducted. Lake Großer Ostersee was fed by alkaline groundwater with a catchment area comprising 35 km^2 (Gruenert and Raeder, 2014).

Sampling

Samples for volumetric shell measurements were taken in June and August 2016 from the epilimnetic layer (0-7 m depth) at the deepest point of each lake. Integrated water samples were taken with a PE sampling tube (7 m length and 4.5 cm diameter) with a 2.5 kg weight at the bottom end. The tube was slowly let down vertically into the water. Its content was filled into a 6 L canister and homogenised by shaking prior to further processing. Samples for the measurements of the *Phacotus* shell abundance and total epilimnetic particulate calcite were

Tab. 1. Characteristics of selected hard water lakes and geographic sampling positions in Bavaria, Germany.

	Gr. Ostersee	Abtsdorfersee	Igelsbachsee	Hopfensee	Altmühlsee
Trophic state	Oligotrophic	Mesotrophic	Mesotrophic	Eutrophic	Hypertrophic
Circulation type	Dimictic	Dimictic	Polymictic	Dimictic	Polymictic
Surface area (ha)	118	78	72	186	450
Max. depth (m)	29.7	20.0	11.5	10.4	2.2
Coordinates (WGS 84)	47°47'25.2'' 011°18'06.5''	47°54'35.4'' 012°54'22.4''	49°08'45.3'' 010°54'13.3''	47°36'06.1'' 010°40'46.6''	49°08'35.8'' 010°43'34.4''
Water temp. (°C)	17.4 (4.7)	15.5 (5.2)	20.9 (2.7)	17.3 (2.8)	22.1 (1.6)
pH value	8.2 (0.2)	7.9 (0.4)	8.6 (0.5)	8.2 (0.4)	9.0 (0.3)
Ca^{2+} conc. (mg L^{-1})	69.2	78.2	37.6	69.3	47.1
TP ($\mu\text{g L}^{-1}$)	>8.2	>17.4	>25*	>35**	>200*
Secci depth (m)	3.3 (0.6)	1.1 (0.2)	3.0 (0.6)	1.1 (0.3)	0.5 (0.2)

Physical, chemical average values of the surface layer (0-7 m) during June till August 2016; TP data from Bavarian Environmental Agency; *2014; **2013; Ca^{2+} concentration from last sampling in August; SD in brackets.

taken between 0-5 m depth to ensure comparability with previous works. Sampling was carried out weekly at the deepest point of each lake during the *Phacotus* growing season from June to October 2015. These samples were taken with a standard Ruttner water sampler and integrated by mixing aliquots inside a measuring cup. Water temperature, pH depth profiles and concentrations of Ca^{2+} in solution were measured according to Gruenert and Raeder (2014).

Infrared gas analysis for carbonate determination

Particulate CaCO_3 concentrations were measured by infrared (IR) CO_2 analysis. We used a revised method with a modified measurement setup (Fig. 1) based on the former conductometric CO_2 analysis reported by Proft (1983, 1984). This enabled a more precise determination of the suspended calcite in lake water samples.

Each lake water sample (200 mL) was filtrated through $0.45 \mu\text{m}$ cellulose-nitrate filters. The filters were oven dried overnight at 50°C and stored in a desiccator before introducing them with tweezers into a reaction vessel containing 10% HCl solution. A carrier gas was streamed through the reaction vessel and absorbed the

emerging CO_2 released by carbonate dissolution. Prior to CO_2 gas analysis with a Saxon Junkalor GmbH (Dessau, Germany) Infralyt 50 gas analyser, the carrier gas was passed through Cu flakes to remove any HCl and dried by passing it through granulated CaCl_2 . The carrier gas was air from the lab, which was pumped with a constant gas flow of 33 mL s^{-1} through NaOH pellets before use to remove the atmospheric CO_2 .

The system was calibrated for a measuring range between $0.005\text{-}0.05 \text{ [mg C L}^{-1}\text{]}$ with a IC-standard solution (containing NaHCO_3 and Na_2CO_3). The Infralyt 50 measured the CO_2 content displayed as ppm CO_2 and the LC-Net-Box transformed the analogue signal to a digital signal for processing with the Jasco (Easton, Maryland, USA) Borwin 1.5 software. The output dimension was given as carbon concentration $c_c \text{ [mg C L}^{-1}\text{]}$. Before each measurement, the background CO_2 (≤ 1 ppm) was set to zero. Measurement time for one sample was 180 s. The 10% HCl solution in the reaction vessel was renewed after the measurement of 10 samples. The conversion of c_c to c_{CaCO_3} concentration was calculated by multiplication of c_c with the ratio of the molecular weights $M_{\text{CaCO}_3} 100.09 \text{ g mol}^{-1}$ and $M_c 12.01 \text{ g mol}^{-1}$ according to equation 1. After every 20 measurements, a

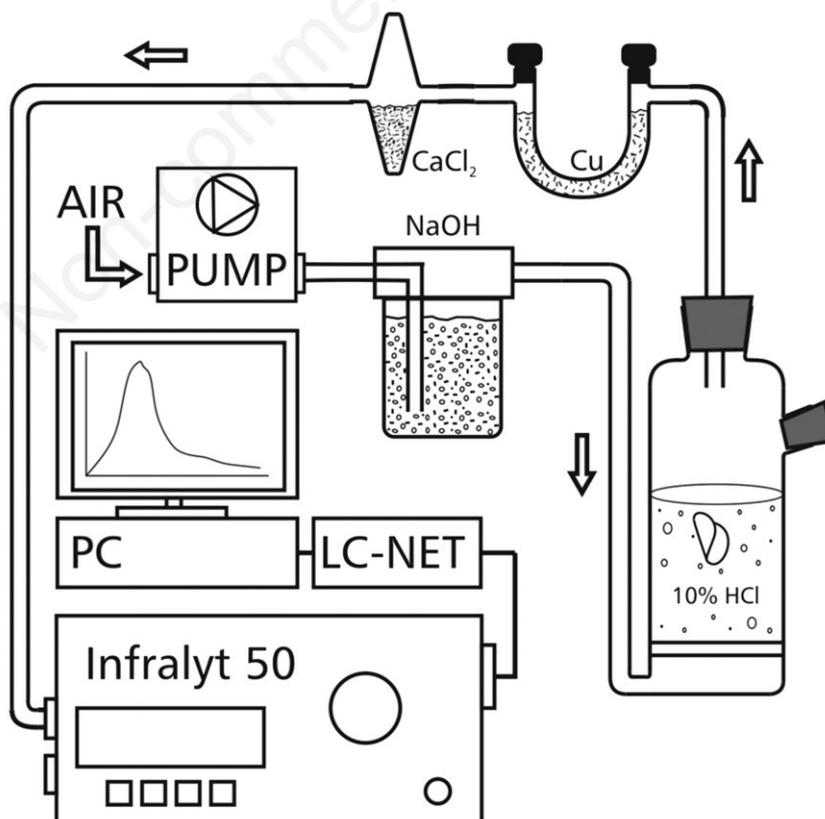


Fig. 1. Schematic representation of the carbonate infrared gas analysis measurement setup.

calibration sample was measured to guarantee accurate results. The system was calibrated after every 60 measurements.

$$c_{CaCO_3}[mg\ L^{-1}] = c_c[mg\ L^{-1}] \times \frac{100.09\ g\ mol^{-1}}{12.01\ g\ mol^{-1}} \quad \text{eq. 1}$$

Plankton analysis

Plankton samples were stored in 100 mL brown glass bottles and preserved with Lugol's solution. Suspended shell concentrations in the water column were determined with an inverted microscope Leitz Labovert at 200-400 \times according to Utermöhl (1959). We counted single shells as one, and complete *Phacotus* housings as two shells. The cell density was derived by dividing the sum of the shells by two because a complete *Phacotus* housing consists of two shells. At least 300 individuals were counted per sample to ensure accurate determination of the shell numbers. Additionally, the shell diameter was determined by taking two perpendicular measurements. From each lake, 150 individuals (50 from three sampling events at different stages of population growth in June, July, and August 2016) were measured with the EOS-Metrology software. To test for mean shell diameter differences between lakes and month of sampling, a two-way-analysis-of-variance (ANOVA) was conducted. In case of significance, Tukey's *post-hoc* tests were computed. Statistical analyses were carried out using the open source software R 3.2.2 (R Core Team, 2015).

Focused ion beam processing

Thin layer cross-sections of a *Phacotus* calcite shell were cut with a FEI (Hillsboro, Oregon, USA) Helios NanoLab G3 UC FIB. Prior to milling, samples were prepared on Si wafers and coated with 20 nm carbon using a Leica EMACE600-Coater. To assure clean cut surfaces, the cutting area in the middle of each shell was sealed with a protective layer of Pt (Fig. 2 b-d) with a thickness of 1 μm , derived from ion induced deposition. The calcite shell was cut and polished using ion currents from 47 nA to 2.5 nA, at an acceleration voltage (AV) of 30 kV. Scanning electron micrographs were obtained at 1.0 kV AV with a viewing angle of 52 $^\circ$ to the ion beam. In total, 23 individuals were cut vertically in the middle and one shell horizontally, parallel to its flat side.

Volume and mass determination

A total number of 24 individuals from five lakes were FIB milled to obtain a site-specific view of the shell morphology and structure. On the shell cross-section images, profile lines along the outer margin of the shell were marked. Two profiles were drawn on each shell

cross-section (Fig. 2d), resulting in 48 profiles of shell halves for cross-section image analysis. On these profiles, the following characteristic parameters (Fig. 2e) were measured (precision as relative error): minimal shell thickness t_{\min} (11.1%), rim width t_{rim} (2.0%), radius r (0.6%), diameter d and height h (1.4%). For each shell, profiles were converted to surface area and rotated 360 $^\circ$ to create a closed up 3D-body of rotation from which the volume (μm^3) was determined by using the CAD software Rhinoceros (McNeel, 2013).

For the estimation of total porosity Φ (%), image analysis was conducted on 15 consecutive cross-section micrographs of two *Phacotus* individuals with the image analysis software FiJi ImageJ (Rasband, 2016). The total porosity was determined in terms of percentage of open pore area per unit shell area on the plane shell cut faces. To eliminate scale errors, we analysed with different sized analysis windows (0.8 \times 0.8 μm and 2.3 \times 0.7 μm) and repeated the measurement on at least five micrographs in series, of which each represented a 0.04 μm deeper cut face of the investigated shell. Subsequently, the shell density ρ (ng μm^{-3}) was calculated according to equation 2 with a density of $\rho_0=0.0027$ ng μm^{-3} for pure calcite.

$$\rho = \rho_0(1 - \Phi) \quad \text{eq. 2}$$

The mass m of CaCO_3 (ng) deposited in each individual *Phacotus* shell was calculated according to equation 3 by multiplying the volume of the 3D-body of rotation with ρ .

$$m = V \cdot \rho \quad \text{eq. 3}$$

RESULTS

Calculation of shell mass

Our measurements of 24 individuals from 5 lakes led to a mean volume of 334.1 μm^3 (SD=70) for a complete *Phacotus* housing with a 13.9 μm diameter (consisting of two shells). Our data showed that shells with the same diameter had a variation in volume up to a factor 1.8. The calculated mean shell mass for a complete specimen (comprising two valves) was 0.86 ng (SD=0.18). The mean shell mass ranged from 1.03 (SD=0.11) ng in lake Großer Ostersee to 0.66 (SD=0.10) ng in lake Igelsbachsee (Tab. 2).

Estimation of porosity

Based on image analysis of the shell cut faces of the cross-section micrographs, highly massive calcite shells were identified (*data not shown*). We decided to calculate the material density according to equation 2 using total porosity derived from image analysis. The cut faces appeared even and showed only few cut pores. We

analysed both a relatively porous and a highly massive appearing shell. The resulting estimated porosities were 2.9% for the porous and 2.2% for the massive shell. The smallest analysed pores had a pore-throat size of 50 nm. Nevertheless, close-up scanning electron micrographs also showed mesopores with diameters smaller than 50

nm (Fig. 3a). These pores certainly contributed to an increased inner surface area but might not contribute to a significantly higher porosity. However, for further calculations, we assumed a total material porosity of $\phi=5\%$ to take this fact into account. According to equation 2, the shell density was $\rho=0.0026 \text{ g } \mu\text{m}^{-3}$.

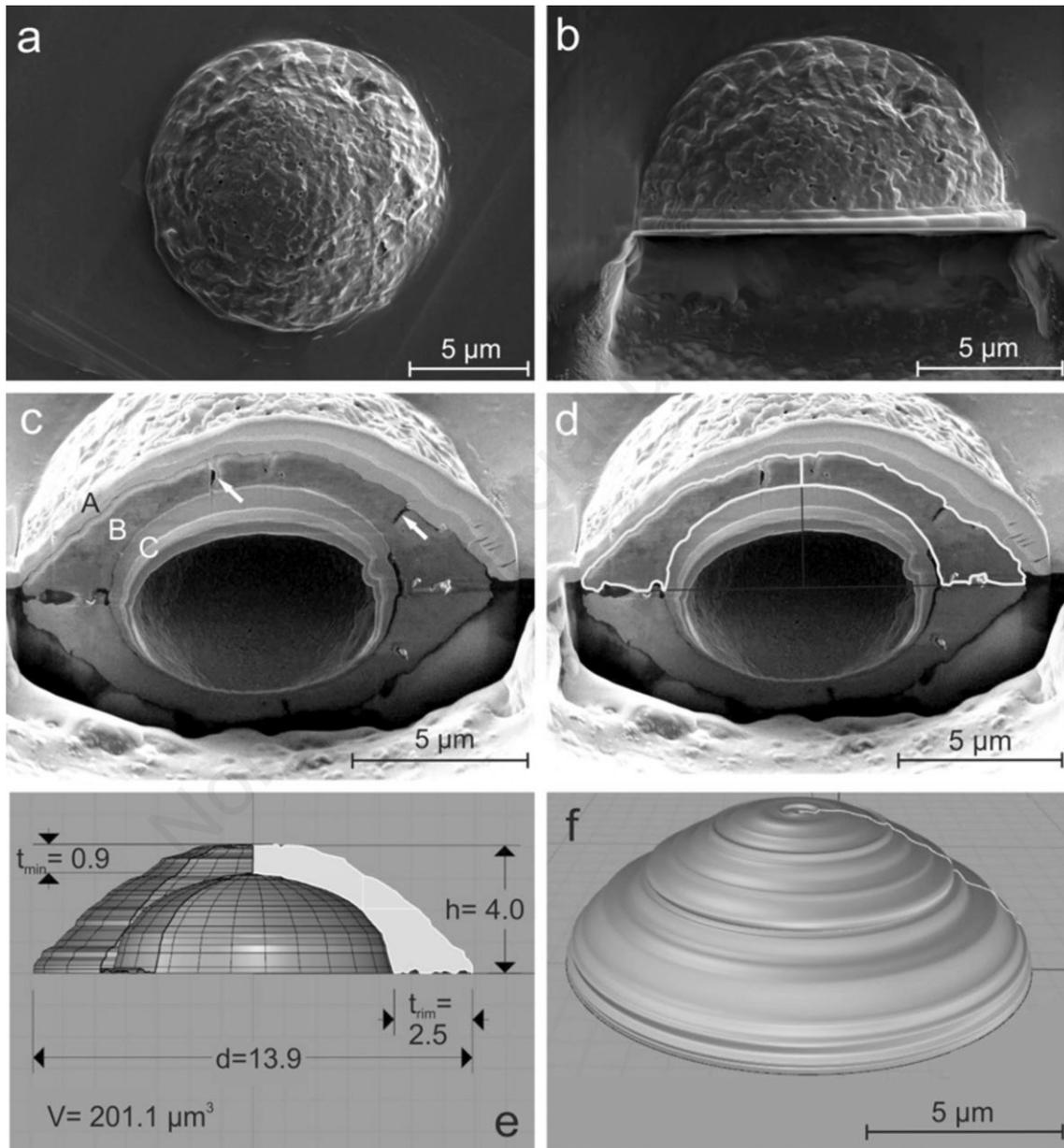


Fig. 2. Method for volume determination, scanning electron micrographs of a *Phacotus lenticularis* shell from lake Großer Ostersee, 4 m depth, June 2015: a) intact shell with flagella pore at the upper right and several small pores on its surface, plan view prior to FIB milling; b) shell, cut in the middle with narrow Pt-protection layer on cutting edge, plan view; c) imaging face of shell cross-section with two exposed pore canals (white arrows), light grey Pt coating (A) over dark grey shell material (B), layers of cutting debris, accumulated inside the shell during cutting process (C), side view; d) shell cross-section with both marked profile lines for creation of a 3D-body, side view; e) open 3D-body of rotation with profile of the cross-section and measuring bars, dimension are in μm , side view perpendicular to profile; f) closed 3D-body for volume determination with Rhinoceros 3D software.

Variability of general dimensions on shell cross-sections

The shell diameter (d) was significantly influenced by the lake's origin ($P=0.007$), but the month of sampling had no significant influence ($P=0.190$). The investigated shells in 2016 had a mean diameter of $13.8 \mu\text{m}$ for lake Abtsdorfersee, $13.1 \mu\text{m}$ for lake Altmühlsee, $12.7 \mu\text{m}$ for

lake Großer Ostersee, $14.0 \mu\text{m}$ for lake Hopfensee, and $12.0 \mu\text{m}$ for lake Igelsbachsee (Fig. 4). The biggest shell was found in lake Abtsdorfersee ($d=21.9 \mu\text{m}$), the smallest shell in lake Igelsbachsee ($d=8.3 \mu\text{m}$).

The shell size parameter height (h) varied significantly between the lakes. The shell thickness (t_{min}) and rim width (t_{rim}) showed significant variation only for the shells from lake Großer Ostersee (Tab. 2). The h/d ratios of the

Tab. 2. General dimensions, volumes and calculated masses of *Phacotus* shells.

	Gr. Ostersee	Abtsdorfersee	Igelsbachsee	Hopfensee	Altmühlsee
Mass* (ng)	0.52 (0.06)	0.43 (0.09)	0.33 (0.05)	0.40 (0.06)	0.48 (0.05)
Volume (μm^3)	201.7 (32.6)	167.0 (36.9)	128.4 (21.4)	155.4 (22.8)	186.7 (21.5)
Diameter (μm)	13.6 (0.9)	14.0 (1.2)	12.5 (0.5)	15.0 (1.2)	15.0 (0.4)
Rim width (μm)	2.4 (0.4)	2.1 (0.4)	1.8 (0.2)	1.6 (0.3)	2.0 (0.3)
Thickness (μm)	1.0 (0.1)	0.7 (0.2)	0.8 (0.2)	0.7 (0.1)	0.8 (0.1)
Height (μm)	4.2 (0.3)	4.5 (0.4)	4.2 (0.1)	4.8 (0.6)	4.8 (0.2)

*Weight for a single shell half with 5% porosity; 48 shell profiles of 24 individuals were measured. Samples were taken in June and August 2016.

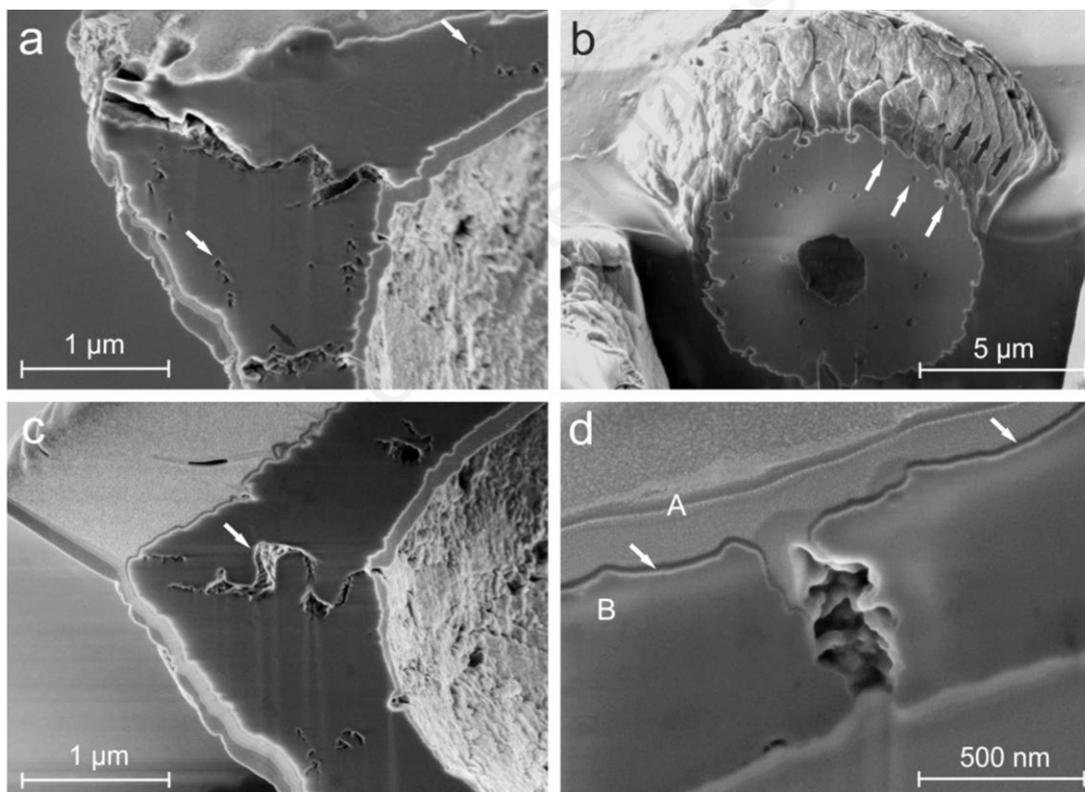


Fig. 3. Close-up scanning electron micrographs of internal pores and cavities on the shell cross-sections: a) contact zone at the shell margin, cut tabular rhombohedral calcite crystals show a step-like cut edge, there are micro pores (white arrows) and a pore-canal (black arrow) in the massive calcite shell; b) imaging face in upper shell zone, horizontally cut parallel to contact planes, tabular rhombohedral calcite crystals (black arrows) are visible on the outer most two rings at the shell margin, inner calcite seems to be very massive and interspersed with few, small pores (white arrows), side view; c) contact zone on the shell rim, calcite crystals are interlocked, the cavity between the two shells (white arrow) formed during imaging process, previously filled with an unknown substance volatile in vacuum under an electron beam; d) shell cross-section in the central zone, pore-canal that intersects the complete shell from inside to outside, its cavernous rim appeared during the imaging process and was also filled with the unknown substance, Pt-protection coating (A), dark 20 nm C-sputter-layer (white arrows), actual shell calcite (B).

Phacotus shells from each lake remained constant (0.32; SD=0.02). The characteristic appearance of the shells from different lakes might be better reflected by the ratio of the shell thickness to its diameter. Shells from two lakes showed remarkable characteristics: the shells from lake Hopfensee had a filigree shell architecture with a low t_{min}/d ratio of 0.045 (SD=0.010), whereas shells from lake Großer Ostersee showed a massive architecture with a high t_{min}/d ratio of 0.075 (SD=0.011). In contrast, the t_{rim}/d ratio was not a good descriptive value because it was dominated by the variation in the shell diameter.

Observations of the shell ultrastructure

The analyses of calcite shell cross-sections from SEM images confirmed that the internal structure consisted of closely stacked tabular calcite crystals. They had approximate dimensions of $1.0 \times 1.6 \times 0.2 \mu\text{m}$. Most of the interspaces were predominately cemented with super-fine-grained calcite lacking a specific crystal structure. Some interspaces were unfilled and formed cylindrical elongated pores that did not penetrate the entire shell. Most pores were not interlinked and ceased before reaching the outer layer. In the inner most shell areas, small pores were pitched in a circular-spiral arrangement along the smaller lateral edge of the tabular crystals (Figs. 2a and 3b). The majority of pores that could be seen at the shell surface were from this type and had an average diameter of $0.11 \mu\text{m}$ (SD=0.03).

Another kind of pore, fewer and likely situated in the central shell area, intersected the whole shell (Figs. 2c and 3a, d). These pore canals had wider pore-throats of approximately $0.2 \mu\text{m}$ and irregular pore walls. According to the behaviour during SEM imaging, these pores seemed to be coated or filled with a volatile organic substance that disappeared under the electron beam under vacuum conditions. The space in the contact zone between the two shells (rim interspace) was filled with the same substance and showed tabular crystals almost interlocked with each other (Fig. 3c).

Phacotus induced calcite precipitation

To quantify the autochthonous calcite precipitation in the epilimnion of lake Großer Ostersee, we measured the amount of total calcite in the upper 5 m of the water column. Remarkably high concentrations of total suspended carbonate of over 1.5 mg L^{-1} were reached from mid-July until mid-September, with a maximum of 3.2 mg L^{-1} (Fig. 5).

During the peak development of *Phacotus lenticularis* on 16th June 2015, the maximum cell density reached $477,800 \text{ ind. L}^{-1}$ in 2 m depth. The average value for the epilimnion (0-5 m) was $153,327 \text{ ind. L}^{-1}$. During this occasion, 21.5% of precipitated calcite was provided by *Phacotus lenticularis*. This peak was followed by a sudden decrease in cell numbers coinciding with rain- and wind-induced epilimnetic mixing and grazing by

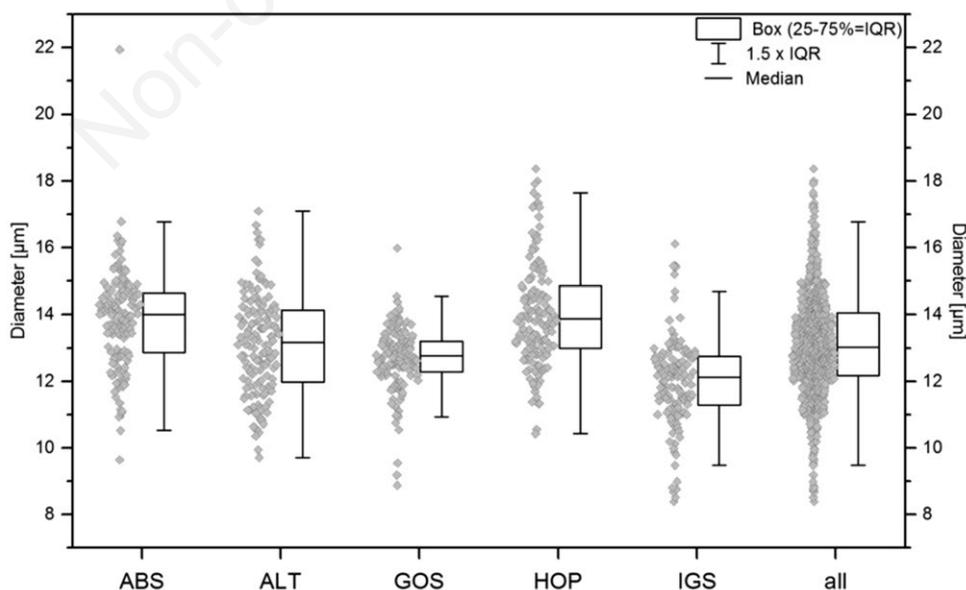


Fig. 4. Variation of *Phacotus* shell diameters measured during cell counting 2016: total n was 150 for each lake; 50 from June, July and August each. Data points as grey squares; summary as box including first to third quartile (25-75%=interquartile range IQR), whiskers describe $1.5 \times \text{IQR}$ (12.5 - 87.5%). The line represents the median. Abbreviations as indicated in methods section.

zooplankton. Three additional *Phacotus* peaks occurred in early July, in late July, and in mid-September. The average amount of precipitated calcite provided by *Phacotus lenticularis* over the whole monitoring period was 6.0% in relation to total suspended calcite owing to the high total precipitated calcite concentrations in the epilimnion.

In the epilimnion of lake Großer Ostersee the average carbonate concentration per square metre of lake surface was $6.53 \text{ g CaCO}_3 \text{ m}^{-2}$ (as average over the whole investigation period from 7th June to 3rd November 2015 in 0-5 m water depth). At peak concentrations on 11th August 2015, the values reached $16.1 \text{ g CaCO}_3 \text{ m}^{-2}$. The total contribution of *Phacotus lenticularis* (6%) on the mean value was $0.38 \text{ g CaCO}_3 \text{ m}^{-2}$ (0-5 m) and reached $0.80 \text{ g CaCO}_3 \text{ m}^{-2}$ during *Phacotus* peak concentrations (21.5%) on 16th June 2015.

DISCUSSION

This study provides a novel methodological basis for a quantitative assessment of *Phacotus* precipitated carbonate in relation to the total carbonate precipitation in lakes. We examined the variations of the *Phacotus*

shape and determined the resulting weight of its shells, which was the most relevant variable in this investigation. Our investigation revealed that the lake's origin significantly influenced the shell diameters and height. Accordingly, the shell volumes and masses varied strongly between lakes. It seems unlikely that dissolution was the reason for the variation in diameter or mass of the shells because neither the water chemistry (Tab. 1), nor the selective electron micrographs of the epilimnetic material (Figs. 2 and 3), showed evidence of dissolution of *Phacotus* shells or calcite crystals in any of the lakes. The shells were found to be highly massive with few pores of different types (*i.e.*, porosity <5%). This resulted in a shell mass for *Phacotus lenticularis* which was 2.8-fold higher than the current estimation of 0.3 ng CaCO_3 per individual reported by Koschel *et al.* (1987).

There might be several reasons that explain why our size-adjusted mass estimation based on 24 individuals from 5 lakes show a higher and more realistic shell mass (0.86 ng CaCO_3) compared to current estimates. First, differences in the methodological approaches need to be considered: in contrast to our study, Koschel *et al.* (1987) had the favourable and rare circumstance of lake water samples in which 100% of the suspended carbonate was derived from *P. lenticularis* cells. Thus, the amount of

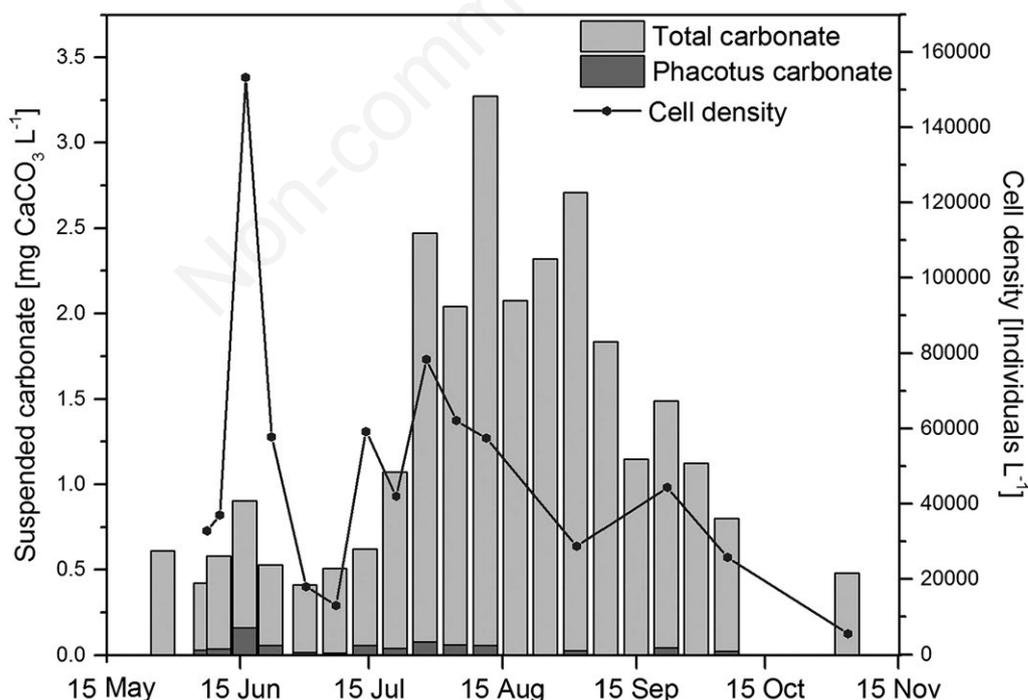


Fig. 5. Total particulate carbonate (light bars) and amount of precipitated calcite provided by *Phacotus lenticularis* (dark bars). Total carbonate values were measured by infrared gas-analysis whereas *Phacotus* carbonate values were calculated using an individual's mass of 1.03 ng (two shells). Not all sampling dates have *Phacotus* cell density data (line). All shown data are integrated for 0-5 m of lake Großer Ostersee in 2015.

carbonate from a certain volume of lake water (on a filter) could be measured conductometrically. The individual shell mass was then calculated through division with the microscopically determined cell concentration.

This approach might have resulted in an inferior shell mass value either because of common inaccuracies during microscopic cell counting or, owing to insufficient transfer of shells from the lake water onto the membrane filters. Homogenisation and division are difficult owing to fairly swift sedimentation of the *Phacotus* shells and can lead to depleted *Phacotus* shell concentrations in decanted water samples.

In our opinion, the methodology we used for volume determination can be considered less prone to errors. Using state-of-the-art FIB technology, the accuracy in imaging was high and scaling for the 3D-software had an absolute precision of less than 0.06 μm and the subsequent volume calculation had an extremely low mean error of $1.3 \times 10^{-5} \mu\text{m}^3$ ($\text{SD}=1.2 \times 10^{-5}$). However, our method for calculating the mass is not entirely unproblematic because the proportion of mesoporosity had to be estimated (in our case 2.4% additional to the measured macroporosity with an average of 2.6%). This resulted in a presumed total porosity of approximately 5%. Accordingly, our calculated shell density ($\rho=0.0026 \text{ ng } \mu\text{m}^{-3}$), was only slightly lower than the of massive calcite. This value is in accordance with the density of biomineralised calcite of planktonic marine algae (dried coccolith powder) measured with a gas pycnometer (Hariskos *et al.*, 2016) and seems plausible in regard to the very massive looking shell cross-sections on the close-up scanning electron micrographs (Fig. 3). In accordance with this, values from Koschel *et al.* (1987) do not seem to be realistic: applying their reported shell density of $\rho_s=0.0006 \text{ ng } \mu\text{m}^{-3}$ to equation 2, the resulting porosity Φ of 77.8% seems exorbitantly high.

Secondly, in the different study lakes the natural variation of the general *Phacotus* size also caused large differences in the mean shell masses and could be an explanation for the difference in shell mass results. In previous publications, these natural variations were not subject to special interest. However, they are highly relevant if the quantitative *Phacotus* contribution to the total carbonate precipitation in lakes is being determined. Our measurements on individual cells from five lakes of different trophic states and mixis showed that *Phacotus lenticularis* had a great variability in size and morphology. Also Koschel and Raidt (1988) identified a strong variation in shell diameter in all of their investigated water bodies. Our investigated shells with diameters in the range of $14.0 \pm 2.4 \mu\text{m}$ are typical. According to Koschel and Raidt (1988) shell diameters over $14 \mu\text{m}$ can be considered as large and Giering (1990) states that shells with smaller diameters (7-12 μm)

“may indicate daughter cells, in which shell development is incomplete”.

Our analyses of the internal shell ultrastructure revealed that the small pores from the crystal interspaces were pitched in a circular-spiral arrangement along the smaller lateral edge of the tabular crystals. This means that we can confirm the oval-circular arrangement of the crystals in the shell centre (Fig. 3b) and the observations of Kampner (1950) as well as those by Koschel and Raidt (1988). They described the rotational symmetry of the crystals at the marginal shell rim and dissymmetrical orientation in the centre of the cell, similar to a “skiodromen model”. Predominantly located in the central shell area, we found a second type of pore, the pore canals. In contrast to the descriptions of Giering (1990), we could not confirm that all of the pores on the inner shell surface were “openings of the pore canals penetrating the *lorica*”. Actually, only a few of them intersected the whole shell from the inside to the outside (Figs. 2c and 3a, d). These pore canals were wider than the pores in the crystal interspaces and showed irregular pore walls.

Furthermore, we can report that shells of vegetative cells do actually touch each other in the contact zone (Fig. 3a) and are sometimes even almost interlocked (Fig. 3c). The rim interspace seemed to be filled with a volatile substance, which under an electron beam under vacuum conditions slowly disappears. This substance might represent the mucilaginous sheath (Hepperle and Krienitz, 1996), previously called “gelatinous mother cell wall” (Giering *et al.*, 1990). We observed that the pore channels seem to be coated or filled with this same volatile organic substance. In our opinion, it seems more probable that the exchange of substances during the motile cell state of *Phacotus lenticularis* is carried out through the pore canals rather than through the rim interspace in the contact zone, as suggested by Kamptner (1950).

Finally, we can report a remarkable contribution of *Phacotus lenticularis* to the total suspended calcite precipitation in lake Großer Ostersee of 21.5% at peak cell densities. Kienitz *et al.* (1993) reported the mean percentages of *Phacotus* calcite of the total suspended epilimnetic calcite in several hard-water lakes of northern Germany ranged from 5%-20% and reached 54% in exceptional circumstances. Fluctuating *Phacotus* populations and irregularly oscillating total calcite precipitation rates were mentioned as reasons for the wide range of variation. Furthermore, the trophic state of the lake has to be considered as an important influencing factor. Significant *Phacotus* calcite percentages are expected predominantly in eutrophic and highly eutrophic lakes (Schlegel *et al.*, 1998).

The total contribution of *Phacotus lenticularis* on the epilimnic mean carbonate concentration per square meter reached up to 21.5% during peak concentrations and on

average 6% over the whole investigation period (7th June - 3rd November 2015). Taking into account that sedimentation is only responsible for about 20% of total current C-turnover from inland waters (Tranvik *et al.*, 2009), and that at least half of the C is precipitated in the form of organic carbon compounds (Noges *et al.*, 2016), carbonate precipitation by *Phacotus lenticularis* represents a small but notable part of C-turnover in lakes. In hard-water lakes, it should certainly be considered in the assessment of lake carbon cycling.

CONCLUSIONS

An accurate quantification of the carbonate sequestration of *Phacotus lenticularis* in lakes requires precise estimation of its shell sizes, masses, and the variations of both variables in relation to spatio-temporal variability. The mean diameter from times during highest cell densities should be used as the basis for all calcite mass calculations. During these events, *Phacotus lenticularis* actually provides relevant amounts of epilimnetic suspended calcite. Further investigation is necessary with respect to persistence, and actual amount of *Phacotus* carbonate stored in recent lacustrine sediments and whether shells tend to be re-dissolved on their way from the epilimnion to the lake bottom.

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REFERENCES

- Cole JJ, Prairie YT, Caraco NF, McDowell WH, Tranvik LJ, Striegl RG, Duarte CM, Kortelainen P, Downing JA, Middelburg JJ, Melack J, 2007. Plumbing the global carbon cycle: Integrating inland waters into the terrestrial carbon budget. *Ecosystems* 10:172-185.
- Giering B, Krienitz L, Casper SJ, Peschke T, Raidt H, 1990. LM and SEM observations on the asexual reproduction and lorica formation of *Phacotus lendneri* Chodat (Chlamydomonadales, Phacotaceae). *Arch. Protistenkd.* 138:75-88.
- Gruenert U, Raeder U, 2014. Growth responses of the calcite-loricated freshwater phytoflagellate *Phacotus lenticularis* (Chlorophyta) to the CaCO₃ saturation state and meteorological changes. *J. Plankton Res.* 36:630-640.
- Hariskos I, Chairpoulou MA, Posten C, Teipel U, Vučak M, 2016. Characterisation of biogenic calcite particles from microalgae-coccoliths as a potential raw material for industrial application. *Ecol. Eng. Environ. Prot.* 7:36-41.
- Hepperle D, Krienitz L, 1996. The extracellular calcification of zoospores of *Phacotus lenticularis* (Chlorophyta, Chlamydomonadales). *Eur. J. Phycol.* 31:11-21.
- Kamptner E, 1950. [Über den Aufbau des Kalkgehäuses von *Phacotus Lendneri* CHOD.]. [Article in German]. *Österr. Bot. Z.* 97: 391-402.
- Koschel R, Proft G, Raidt H, 1987. [*Phacotus*-Massenentwicklungen - eine Quelle des autochthonen Kalkeintrages in Seen]. [Article in German]. *Limnologica* 18:457-459.
- Koschel R, Raidt H, 1988. [Morphologische Merkmale der *Phacotus*-Hüllen in Hartwasserseen der Mecklenburger Seenplatte]. [Article in German]. *Limnologica* 19:13-25.
- Krienitz L, Koschel R, Giering B, Casper SJ, Hepperle D, 1993. Phenomenology of organismic calcite precipitation by *Phacotus* in hardwater lakes and ponds of northeastern Germany. *Verh. Int. Verein. Limnol.* 25:170-174.
- McNeel R, 2013. Rhinoceros: CAD software, ver. 5.1.30129.1756. Robert McNeel & Associates, Barcelona.
- Müller G, Oti M, 1981. The occurrence of calcified planktonic green algae in freshwater carbonates. *Sedimentology* 28:897-902.
- Noges P, Cremona F, Laas A, Martma T, Room E-I, Toming K, Viik M, Vilbaste S, Noges T, 2016. Role of a productive lake in carbon sequestration within a calcareous catchment. *Sci. Total Environ.* 550:225-230.
- Pocratsky LA, 1982. Nutritional, chemical and ultrastructural characterization of the lorica and extracellular mucilage of *Phacotus lenticularis* (Phacotaceae, Volvocales). PhD Thesis, University Tennessee.
- Proft G, 1983. [Automatische Bestimmung von Kohlendioxid und Carbonat in Wasser nach dem flow-stream-Prinzip]. [Article in German]. *Acta Hydroch. Hydrob.* 11:235-239.
- Proft G, 1984. [Die pelagische Calcitfällung und der Carbonatgehalt von Sedimenten pleistozäner Seen]. [Article in German]. *Acta Hydroch. Hydrob.* 12:321-326.
- R Core Team, 2015. R: A Language and Environment for Statistical Computing, ver. 3.2.2. R Foundation for Statistical Computing, Vienna.
- Rasband WS, 2016. ImageJ: public domain Java image processing program, ver. v1.51h. National Institutes of Health, Bethesda.
- Schlegel I, Koschel R, Krienitz L, 1998. On the occurrence of *Phacotus lenticularis* (Chlorophyta) in lakes of different trophic state. *Hydrobiologia* 369-370:353-361.
- Schlegel I, Krienitz L, Hepperle D, 2000. Variability of calcification of *Phacotus lenticularis* (Chlorophyta, Chlamydomonadales) in nature and culture. *Phycologia* 39:318-322.
- Siemińska J, 1952. The plankton of the artificial lake at the Roznow dam. *Mem. Acad. Pol. Sci. B* 18:1-109.
- Sladeczek V, Cyrus Z, Borovickova A, 1958. Hydrobiological investigations of a treatment of beet sugar factory's wastes

- in an experimental lagoon. Sbornik vus. skolu chem. techn. paliv a vodu: 121-150.
- Steinberg C, Klee R, 1983. [Röntgenmikroanalyse der Schale einer einzelligen Grünalge: Chemismus einer Phacotus lundneri-Schale]. [Article in German]. Mikrokosmos 72: 170-173.
- Tranvik LJ, Downing JA, Cotner JB, Loiselle SA, Striegl RG, Ballatore TJ, Dillon P, Finlay K, Fortino K, Knoll LB, Kortelainen PL, Kutser T, Larsen S, Laurion I, Leech DM, McCallister SL, McKnight DM, Melack JM, Overholt E, Porter JA, Prairie Y, Renwick WH, Roland F, Sherman BS, Schindler DW, Sobek S, Tremblay A, Vanni MJ, Verschoor AM, Wachenfeldt E von, Weyhenmeyer GA, 2009. Lakes and reservoirs as regulators of carbon cycling and climate. Limnol. Oceanogr. 54:2298-2314.
- Utermöhl H, 1958. [Zur Vervollkommnung der quantitativen Phytoplankton-Methodik]. [Book in German]. E. Schweizerbart, Stuttgart: 38 pp.

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