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Data Article

Data on euglyphid testate amoeba densities, corresponding protozoic silicon pools, and selected soil parameters of initial and forested biogeosystems



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ABSTRACT

The dataset in the present article provides information on protozoic silicon (Si) pools represented by euglyphid testate amoebae (TA) in soils of initial and forested biogeosystems. Protozoic Si pools were calculated from densities of euglyphid TA shells and corresponding Si contents. The article also includes data on potential annual biosilicification rates of euglyphid TA at the examined sites. Furthermore, data on selected soil parameters (e.g., readily-available Si, soil pH) and site characteristics (e.g., soil groups, climate data) can be found. The data might be interesting for researchers focusing on biological processes in Si cycling in general and euglyphid TA and corresponding protozoic Si pools in particular.

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| Subject area | Biology |
|----------------------------|---|
| More specific subject area | Biogeochemistry |
| Type of data | Tables |
| How data was acquired | Soil analyses (for details see Section 2.2.) and microscopical exam- inations |
| | (for details see Section 2.3.) |
| Data format | Analyzed data |
| Experimental factors | Soil parameters, euglyphid testate amoeba densities, and correspond- ing protozoic silicon pools |
| Experimental features | Analyses of initial (chronosequence) and forest sites |
| Data source location | Germany, for latitudes & longitudes and further site details see Table 1. |
| Data accessibility | All data are presented within the paper. |

Specifications table

Value of the data

- The provided dataset is useful for comparison with the results of other authors regarding, e.g., ecological (euglyphid testate amoeba densities) and biogeochemical (protozoic silicon pools, annual biosilicification) issues in different global ecosystems.
- Data on protozoic silicon pools and corresponding annual biosilicification rates might emphasize the need for detailed investigations of silicon (re-)cycling in unicellular organisms in general and testate amoeba in particular (e.g., qualitative characterization of biogenic silicon, isotope analysis).
- Together with other datasets the presented data allow meta-analyses to examine significant controls (steady state soils) and dynamics (initial soils) of euglyphid testate amoeba densities and related amoebal biosilicification processes in soils in detail.
- In combination with data of other authors the presented data can be used for modelling to assess the role of euglyphid testate amoebae (compared to other organisms that synthesize biogenic silicon, e.g., plants or diatoms) for silicon cycling in soils and corresponding silicon fluxes from terrestrial to aquatic ecosystems.

1. Data

Diverse unicellular and multicellular organisms are able to synthesize structures of amorphous silica $(SiO_2 \cdot nH_2O)$ [1]. In soils these structures represent different biogenic silicon (BSi) pools depending on their origin [2,3]. In general, BSi plays an important role in the global cycling of Si [4–6]. However, while research has been focused on plantal Si and corresponding phytogenic Si pools since decades [e.g., 7–9], far less is known about other BSi pools, e.g., protozoic Si pools represented by euglyphid testate amoeba (TA) shells [3,10–15]. Euglyphid TA represent a monophyletic clade (Euglyphida) of unicellular soil protists with a self-secreted siliceous test (shell) and a worldwide distribution [16].

The presented data were the basis for analyses of protozoic Si pools in initial [12] and forested biogeosystems [3]. The dataset in the present article provides information on i) site characteristics, geographic positions, and climate data of the initial and forested biogeosystems (Table 1), ii) analyzed soil parameters (Table S1), iii) densities of euglyphid TA shells in soils (Table S2), and iv) corresponding protozoic Si pools as well as annual biosilicification rates of euglyphid TA at these sites (Table S3).

Table 1

Site characteristics, geographic positions, and climate data of the examined initial and forested biogeosystems. Data on precipitation and temperature represent annual averages for the period 1981–2010 (German Meteorological Service).

| Site | Lithology | Soil group (WRB) | Humus form | Ecosystem | Coverage (%) | | | Geographic position | | Altitude | Precipitation | Temperature |
|------|------------------------------|---------------------|---------------|---------------------------------------|--------------|-------|------|---------------------|--------------|------------|---------------|-------------|
| | | (WKB) | IoIIII | | Tree | Shrub | Herb | Latitude | Longitude | (m a.s.l.) | (mm) | (°C) |
| сс | Spoil of Quaternary sands | Arenosol | - | Initial (artificial catchment) | - | - | - | 51°36′18″ N | 14°15′58′′ E | 135 | 568 | 9.6 |
| NL | Spoil of Quaternary sands | Arenosol | - | Initial (artificial catchment) | - | - | - | 51°35′50″ N | 14°17′22″ E | 132 | 568 | 9.6 |
| AB | Calcareous glacial till | Stagnosol | Mull | Forest (beech, oak) | 68 | 3 | 70 | 54°12′23″ N | 13°02′21″ E | 34 | 654 | 8.6 |
| EG | Dolomitic limestone | Cambisol | Mull | Forest (beech) | 65 | 0 | 78 | 48°20′19′′ N | 09°25′42″ E | 786 | 808 | 9.5 |
| HE | Basalt | Vertisol | Mull | Forest (beech, ash, spruce, maple) | 65 | 3 | 60 | 47°47′53″ N | 08°45′08″ E | 703 | 820 | 9.2 |
| MR | Calcareous loess | Luvisol | Mull | Forest (beech) | 85 | 35 | 50 | 49°07′13′′ N | 08°40′38″ E | 217 | 776 | 10.0 |
| PP | Calcareous loess | Stagnosol | Mull | Forest (pine, beech) | 70 | 0 | 2 | 48°51′17′′ N | 08°46′14′′ E | 473 | 889 | 9.5 |
| ZE | Kaolinitic claystone | Stagnosol | Mull | Forest (hornbeam, oak) | 85 | 10 | 60 | 48°38'37'' N | 09°33′13′′ E | 390 | 1090 | 9.9 |
| HS | Siliceous sandstone | Podzol | Moder | Forest (pine, spruce) | 73 | 18 | 63 | 48°34′21″ N | 08°20′60′′ E | 832 | 1607 | 8.1 |
| RO | Calcareous glacial till | Luvisol | Moder | Forest (beech, oak) | 70 | 5 | 5 | 53°30′38″ N | 13°21′43″ E | 71 | 530 | 8.7 |
| SL | Eolian sands (dune) | Arenosol | Moder | Forest (pine, beech) | 80 | 0 | 5 | 49°20′51″ N | 08°38′14′′ E | 123 | 861 | 11.0 |
| НК | Siliceous sandstone | Planosol | Peat | Forest (spruce, pine) | 65 | 15 | 60 | 48°34′13″ N | 08°20′53′′ E | 907 | 1607 | 8.1 |

2. Experimental design, materials and methods

2.1. Site description and sampling scheme

2.1.1. Initial sites

The artificial catchments 'Chicken Creek' (CC) and 'Neuer Lugteich' (NL) are part of a post-mining landscape located in the active mining area 'Welzow-South' (lignite open-cast mining, 150 km southeast of Berlin) in the state of Brandenburg, Germany. Climate is characterized by an average air temperature of 9.6 °C with an annual precipitation of 568 mm comprising data from 1981 to 2010.

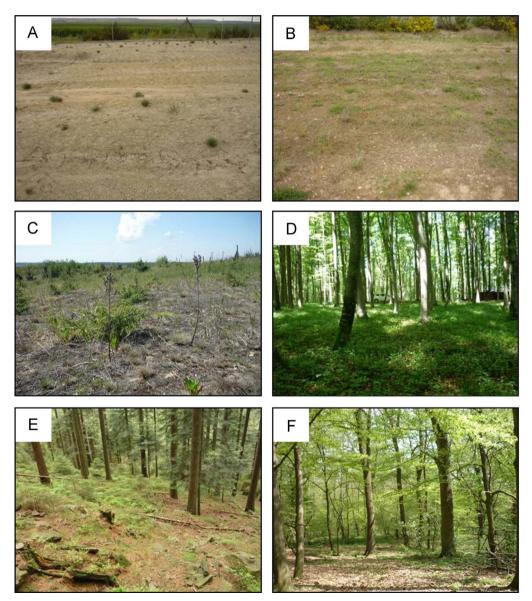


Fig. 1. Photographs of initial (a-c) and some selected forest sites (d-f). A) CC (3-year-old), b) CC (5-year-old), c) NL (10-year-old), d) EG, e) HS, and f) RO. For site details see Table 1 and Section 2.1.

The construction of CC was completed in 2005 (time zero). In 2008 a small area in the west of the catchment was again restored to time zero (removal of the upper 20 cm of soil) for additional experimental plots. Construction of NL was finished in 2001 (time zero). Soils classify as Protic Arenosol (Calcaric, Transportic) or Haplic Arenosol (Hyperochric, Transportic) depending on site age [17]. Detailed information on site construction of CC and NL can be found in Gerwin et al. [18] and Kendzia et al. [19], respectively. All samples were taken from Quaternary substrate at 3-, 5- (CC), and 10-year-old (NL) spots representing a chronosequence (Fig. 1a-c). Samples ($20 \text{ cm} \times 20 \text{ cm} \times 5 \text{ cm}$; sub-divided in two compartments: 0–2.5 and 2.5–5 cm depth) were taken at randomly chosen spots within an area of approx. 25 m^2 . Vegetated (cov) and uncovered (unc) spots were sampled in four field replicates each to analyze possible impacts of vegetation (3 cov: *Tussilago farfara* and *Trifolium arvense*; 5 cov: *Corynephorus canescens* and *T. arvense*) on protozoic Si pools. At NL almost the whole surface was vegetated with biogenic crusts, Poales, and several shrubs, which is why only vegetated spots (10 cov) were sampled. Samples were taken in May 2010 (CC: 5 unc, 5 cov), May 2011 (NL: 10 cov), and August 2011 (CC: 3 unc, 3 cov).

2.1.2. Forest sites

Ten non-eroded forest sites showing huge differences in climate, parent material, and pedogenesis were selected. Mean annual precipitation rates range from 530 to 1600 mm, mean annual air temperatures from 8 to 11 °C. Soils comprise (i) a sandy Arenosol developed from eolian sands (SL, dune), a Podzol and a Planosol from siliceous sandstones (HS, HK) very low in weatherable minerals (< 10% feldspars, mica), (ii) silty to loamy Luvisols and Stagnosols from calcareous, illitic loess (MR, PP) and sandy to loamy Luvisols and Stagnosols from glacial till (RO, AB), both parent materials with intermediate contents of weatherable minerals (feldspars, mica), (iii) a clayey Cambisol from dolomitic limestone (EG), (iv) a clayey Stagnosol from kaolinitic claystone (ZE), and finally (v) a clayey, smectitic Vertisol from basalt (HE) very high in weatherable minerals, like augite and plagioclase. The forest stands are old and are assumed to be in steady state in terms of TA dynamics at a decadal time scale (photographs of some selected forest sites can be found in Fig. 1d-f). Soil samples were taken in four field replicates (n = 4) at all sample sites except HK (n = 3). The field replicates $(20 \text{ cm} \times 20 \text{ cm})$ each) were placed randomly within an area of approx. 100 m². Per field replicate samples were taken in the upper 5 cm (incl. organic layers except for fresh litter) differentiating between two superimposed soil compartments about the same size (ideally 20 cm \times 20 cm \times 2.5 cm each) and transferred to plastic bags. Sampling took place in spring 2010 within six weeks (April 26th–June 6th).

2.2. Soil parameters

Bulk densities (BD, g cm⁻³) were calculated by dividing weights of oven-dried (105 °C) aliquots of soil samples by corresponding volumes. Remaining soil samples were air dried and sieved (2 mm) to separate fine-earth (<2 mm) from skeleton content (>2 mm). For soil analyses only fine-earth was used.

2.2.1. Soil pH, carbon, and nitrogen

Soil pH was measured using a glass electrode in a 0.01 M CaCl₂ solution with a soil-to-solution ratio of one-to-five. For total carbon as well as nitrogen analyses (C_t and N_t) fine-earth samples were finely powdered in a disc mill. Subsequently, C_t and N_t were determined by dry combustion using an elemental analyzer (CNS TruSpec, Leco Instruments). Total inorganic carbon (TIC) was measured with a multiphase analyzer (RC 612, Leco Instruments). Soil organic carbon (SOC) concentrations were calculated by subtraction (C_t -TIC) and C:N ratios were calculated by division (SOC:N_t). Soil C and N analyses were performed at the minimum of two lab repetitions per sample.

2.2.2. Readily-available silicon

For extraction of the calcium chloride $(CaCl_2)$ soluble, so-called readily- or plant-available Si fraction (Si_{CaCl_2}) , 2 g of soil was mixed with 20 ml of a 0.01 MCaCl_2 solution per sample and continuously shaken for 16 h using a lab roller mixer [20]. This Si fraction was extracted to characterize the Si supply for shell synthesis of euglyphid TA in soils. Subsequent to extraction, the extracts were

centrifuged (4000 rpm, 30 min), filtered using 0.45 µm polyamide membrane filters, and Si concentrations were determined by ICP-OES (iCAP 6300 Duo, Thermo Scientific). Complete extraction work was done using plastic equipment only and results represent arithmetic means of three lab repetitions per sample.

2.2.3. Data conversion and calculation steps

All results except for pH were converted to an oven-dry basis (105 °C). Fine-earth mass (FEM in kg m⁻²) was calculated considering bulk density, thickness and skeleton content (wt%). Total FEM (FEM_t) of the upper 5 cm was calculated as the sum of FEM of superimposed compartments. For the upper 5 cm of soil pH was averaged as follows: Per compartment pH was multiplied with the corresponding FEM, divided by FEM_t and subsequently these results were summed up. Mass densities (g m⁻²) of SOC, N_t, and Si_{CaCl2} were calculated compartment-wise by multiplying FEM with element concentrations (g kg⁻¹). Finally, the results of superimposed compartments were summed up for the upper 5 cm of soil.

2.3. Euglyphid testate amoeba densities, protozoic silicon pools, and annual biosilicification

Soil samples in the plastic bags were homogenized by gentle manual mixing and subsequently 2 g of fresh soil was taken per sample for TA analyses and stored in 8 ml of formalin (4%). Soil suspensions received from serial dilution (1000–31.25 mg soil in 8 ml of water each) were stained with aniline blue. TA were enumerated using an inverted microscope (OPTIKA XDS-2, magnifications of 200 × and 400 ×) differentiating between full (living incl. encysted individuals, stained) and empty shells (unstained) of the order Euglyphida. TA densities (shells cm⁻²) were calculated considering TA shell numbers (g^{-1} dry weight), bulk density (g cm⁻³), and thickness (cm) per soil compartment. TA densities of the upper 5 cm were calculated by summing up the corresponding TA densities of superimposed soil compartments.

For calculation of protozoic Si pools we differentiated between different TA taxa of the order Euglyphida with known silica contents per shell as published by Aoki et al. [10]. Summarizing these data we calculated Si contents (pg shell⁻¹, in parentheses listed below) by simple multiplication (SiO₂ content \times 28/60 = Si content) for 9 TA taxa: Assulina muscorum (750), Corythion dubium (580), Euglypha spec. (720), Euglypha rotunda/laevis type (420), Euglypha strigosa type (1420), Tracheleuglypha dentata (750), Trinema complanatum (500), Trinema enchelys (770), and Trinema lineare (360). Indistinctly euglyphid shells or other silica platelet synthesizing TA taxa (e.g., Valkanovia elegans) were recorded as 'euglyphid TA' (700 pg Si per shell, mean of the Si content per shell of the 9 taxa above).

Protozoic Si pools (BSi_{TA} ; mg m⁻²) were calculated per soil compartment using the following formula:

$$BSi_{TA} = \sum_{i=1}^{n} (N_i \times Si_i \times \rho_b \times t \times 10^{-5})$$
(1)

where N_i is the number of euglyphid TA shells (g⁻¹ dry weight), Si_i is the corresponding Si content (pg shell⁻¹; given in parentheses listed above), ρ_b is the bulk density (g cm⁻³), and *t* is the thickness (cm) of the corresponding soil compartment. In contrast, for estimation of annual biosilicification only living euglyphid TA (g⁻¹ dry weight) were considered for N_i in Eq. (1) due to their ability of reproduction. After calculation steps as described in Eq. (1) results were multiplied with 13 and 90 (potential TA generations per year, see Foissner [21]) for minimal and maximal annual biosilicification rates, respectively. For calculation of protozoic Si pools and euglyphid TA biosilicification rates of the upper 5 cm of soil the results of superimposed soil compartments were added up.

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Transparency document. Supplementary material

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Appendix A. Supplementary material

Supplementary data associated with this article can be found in the online version at https://doi.org/10.1016/j.dib.2018.10.164.

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