



HAL
open science

Revisiting Early Jurassic Biscutaceae: *Similiscutum giganteum* sp. nov.

Samuel Mailliot, Emanuela Mattioli, Micaela Chaumeil Rodríguez, Bernard Pittet

► **To cite this version:**

Samuel Mailliot, Emanuela Mattioli, Micaela Chaumeil Rodríguez, Bernard Pittet. Revisiting Early Jurassic Biscutaceae: *Similiscutum giganteum* sp. nov.. *Journal of Micropalaeontology*, 2023, 42 (1), pp.1-12. 10.5194/jm-42-1-2023. hal-04821814

HAL Id: hal-04821814

<https://hal.science/hal-04821814v1>

Submitted on 6 Dec 2024

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution 4.0 International License



Revisiting Early Jurassic Biscutaceae: *Similiscutum giganteum* sp. nov.

Samuel Mailliot¹, Emanuela Mattioli^{1,2}, Micaela Chaumeil Rodríguez^{1,3,4}, and Bernard Pittet^{1,†}

¹Univ Lyon, UCBL, ENSL, UJM, CNRS, LGL-TPE, 69622, Villeurbanne, France

²Institut Universitaire de France (IUF), Paris, France

³División Geología, Museo de La Plata, FCNYM, UNLP, La Plata, Buenos Aires, Argentina

⁴YPF TECNOLOGÍA S.A., Laboratorio de Bioestratigrafía, Buenos Aires, Argentina

†deceased

Correspondence: Emanuela Mattioli (emanuela.mattioli@univ-lyon1.fr)

Received: 13 September 2022 – Revised: 26 January 2023 – Accepted: 6 February 2023 – Published: 3 March 2023

Abstract. A large, broadly elliptical coccolith of the genus *Similiscutum* (Biscutaceae) was observed in sediments dated from the Lower Jurassic (upper Pliensbachian to Toarcian) coming from different localities of western Tethys, namely Portugal (Lusitanian Basin), France (Causses and Paris basins) and Spain (Subbetic area). This form is quite easy to find in the Toarcian GSSP (Global Stratotype Section and Point) of Peniche (Portugal), where the holotype has been described. More than 100 specimens of *Similiscutum* were digitally captured using a CCD camera, including this large form and two other related species, *Similiscutum finchii* and *Similiscutum novum*. The length and width of the coccoliths and the length and width of their central area were measured, and biometric analyses were performed. Results show that this large morphotype of *Similiscutum* is well characterized and easily differentiable by its size and morphology from the species *S. finchii* and *S. novum*, which are characterized by a similar extinction pattern in optical-microscope crossed polars. On the basis of combined differences in size and in central-area shape and structure, *Similiscutum giganteum* sp. nov. is introduced here. (Plant Fossil Names Registry no.: PFN003067; Act LSID: urn:lsid:plantfossilnames.org:act:3067).

1 Introduction

The Biscutaceae are the oldest known placolith coccoliths, appearing at the very base of the Pliensbachian (Bown, 1987a, b; de Kaenel and Bergen, 1993; Mattioli and Erba, 1999; Fraguas et al., 2015, 2018; Peti et al., 2017; Ferreira et al., 2019). During the Pliensbachian and the early Toarcian, this family underwent an important diversification and became a major component of Lower Jurassic calcareous nannofossil assemblages (Bown, 1987a; Mattioli and Erba, 1999; Wiggan et al., 2018).

Lower Jurassic coccoliths belonging to the family Biscutaceae Black, 1971, emend. Bown, 1987a, have been relatively well known since the publication of taxonomic works by Bown (1987a) and de Kaenel and Bergen (1993), where the genus *Similiscutum* was first introduced, and by Mattioli et al. (2004), where the differentiation between Lower

Jurassic species belonging to the genera *Biscutum* and *Similiscutum* was quantitatively assessed using biometrics. However, a large *Similiscutum* species, morphologically close to *Similiscutum finchii* from an ultrastructural and optical point of view, seems to stay undescribed yet. This morphotype was observed in several sections of western Tethys, namely in Portugal (Peniche section; Mailliot, 2006; Ferreira et al., 2019), France (Causses and distal margin of Armorican Massif; Mailliot, 2006; Menini et al., 2019) and Spain (Subbetic area; Reolid et al., 2014). In the literature, Bown (1987a) already mentioned very large specimens of *Similiscutum finchii* recorded in an Argentinian section (Picún Leufú). Subsequently, other papers figured this form to be a representative of *Biscutum* (= *Similiscutum*) *finchii* (e.g. Perilli and Comas-Rengifo, 2002; Perilli and Duarte, 2006; Fraguas et al., 2008; see list of synonymies for further details).

De Kaenel and Bergen (1993) separated the genus *Similiscutum* from *Biscutum* and described several new species in *Similiscutum*; this allowed an improved biochronological subdivision of the Jurassic period. Since then, the first occurrence of placoliths of *Similiscutum* have been used to define the lowermost nannofossil zone of the Pliensbachian stage (Bown and Cooper, 1998; Mattioli and Erba; 1999; Fraguas et al., 2015, 2018; Peti et al., 2017; Ferreira et al., 2019). This event is now used in the Geologic Time Scale (Hesselbo et al., 2020).

Using biometric analysis, this study aims to quantitatively assess the morphology of the large form of *Similiscutum* and to differentiate it from the other related *Similiscutum* species, such as *S. finchii* and *S. novum*. Given its distinctive morphology and its stratigraphic range restricted to upper Pliensbachian and Toarcian strata (Ferreira et al., 2019), the potential stratigraphic use of this new species has to be explored.

2 Material and methods

Two rock samples characterized by high absolute abundances of nannofossils and a fairly good coccolith preservation were selected around the Pliensbachian–Toarcian boundary of Peniche (Ponta do Trovão section) in Portugal (Table 1), where the new *Similiscutum* morphotype is relatively easily recovered compared to other samples or localities. In these two samples, images of a minimum of 15 coccoliths were digitally captured for each of three species – *Similiscutum finchii*, *S. novum* and a large form of *Similiscutum* – in both cross-polarized light and parallel polars. Image acquisition was made using a CCD video camera (Sony XC-77CE) on an “Axioskop 40” Zeiss microscope, using the software “ITI camera configurator” (version 3.3.0.0).

The terminology used in this study follows the guidelines of Young et al. (1997). For each specimen, four simple parameters were analysed: the length and width of the coccoliths were measured on parallel polar images, where the outline is more easily detected; the length and width of the coccolith central area were measured on polarized-light images. Measurements were made using the software “Scion Image” (beta version 4.0.2), which is a modified Windows equivalent of the “NIH Image” technology developed for Macintosh. The accuracy of measurements was estimated by multiple-scale calibration (conversion of pixels into micrometres using “Scion Image”) and by multiple dimensional measurements on the same specimen for the four considered parameters. The scale is $1\ \mu\text{m} = 9.34 \pm 0.02$ pixels; the accuracy of scale calibration is $\pm 0.002\ \mu\text{m}$. Mean error on dimensional measurements is $\pm 0.088\ \mu\text{m}$. By adding these two types of error, the accuracy of dimensional measurements is $\pm 0.09\ \mu\text{m}$, which is very close to the error of $\pm 0.1\ \mu\text{m}$ published by Young et al. (1996) for similar biometric methods.

At first, mixture analysis was applied to coccolith length and width of the entire population of measured specimens

using the PAST 3.01 software (Hammer et al., 2001). The data were then treated with the software “StatView” (version 5.0). Simple parameters were considered for the morphologic description of the three species, such as the coccolith and central-area dimensions (length and width), the relative proportion of the central area with respect to the coccolith size (calculated as the ratios “central-area length over coccolith length” and “central-area width over coccolith width”), and the coccolith ellipticity (calculated as the ratio “coccolith length over coccolith width”). The frequency distribution of these parameters was compared between the considered species in order to test if the large form of *Similiscutum* can be significantly differentiated by its size and shape from *S. finchii* and *S. novum*. According to the work of Mattioli et al. (2004), a 10th–90th percentile interval was chosen to distinguish between a pair of species by their dimensions. This means that two species can be differentiated by means of one of the considered parameters if the frequency distributions of such parameter are different for more than 80 % of the populations (Mattioli et al., 2004, see Fig. 5). In order to test if the average of measured parameters is significantly different between two given species, the ANOVA (ANalysis Of VAriance) post-hoc Bonferroni–Dunn test was applied. This test evaluates the probability (p) of having the same average for a given parameter for a pair of species. In this study, for a significance level at 5 % in a Bonferroni–Dunn test, the significant level of differences between the average values of a pair of species is $p < 0.0167$. The new measurements acquired in this paper were then compared to published biometric data of Lower Jurassic species of Biscutaceae (Mattioli et al., 2004).

3 Results

3.1 Mixture analysis

A comparison of one- and two-group models was performed using the whole dataset for coccolith length and width by comparing their associated Akaike information criterion (AIC; Akaike, 1973) according to the statistical procedure developed by Favre et al. (2008) that allows the optimized separation of two sets of normally distributed values. This method is thus based on two a priori-defined sets of values – namely, one with the smallest mean and another with the largest mean – obtained for a given, normally distributed variable (Suchéras-Marx et al., 2010). The number of size classes or bins was chosen in order to obtain each bin at $0.5\ \mu\text{m}$. The results of mixture analysis show two sets of samples with a mode at $4.0\text{--}5.0$ and $7.5\text{--}8.5\ \mu\text{m}$ for coccolith length and a mode at $3.5\text{--}4.5$ and $6.5\text{--}7.5\ \mu\text{m}$ for coccolith width (Fig. 1). According to mixture analysis, the dataset has a bimodal nature with a larger morphotype (herein named *Similiscutum giganteum* sp. nov.; Plate 1) and two smaller morphotypes, namely *S. finchii* and *S. novum* (Plate 2). In the following, we present boxplot analysis of coccolith geometry separately for the three species.

Table 1. Stratigraphic information of the studied samples. Repository: Collections de Géologie de Lyon.

Sample	Location	Age	No. of measured specimens	Preservation	Absolute abundance of nannofossils per grams of rock
Pen25bis–To2003 FSL no. 766615	Peniche, Portugal	Lower Toarcian (Polymorphum Zone) = Bed 16c (Mouterde, 1955)	54	Good	781×10^6
Pen08–Do2003 FSL no. 766528	Peniche, Portugal	Upper Pliensbachian (Emaciatum Zone) = Bed 15a (Mouterde, 1955)	48	Moderate to good	702×10^6

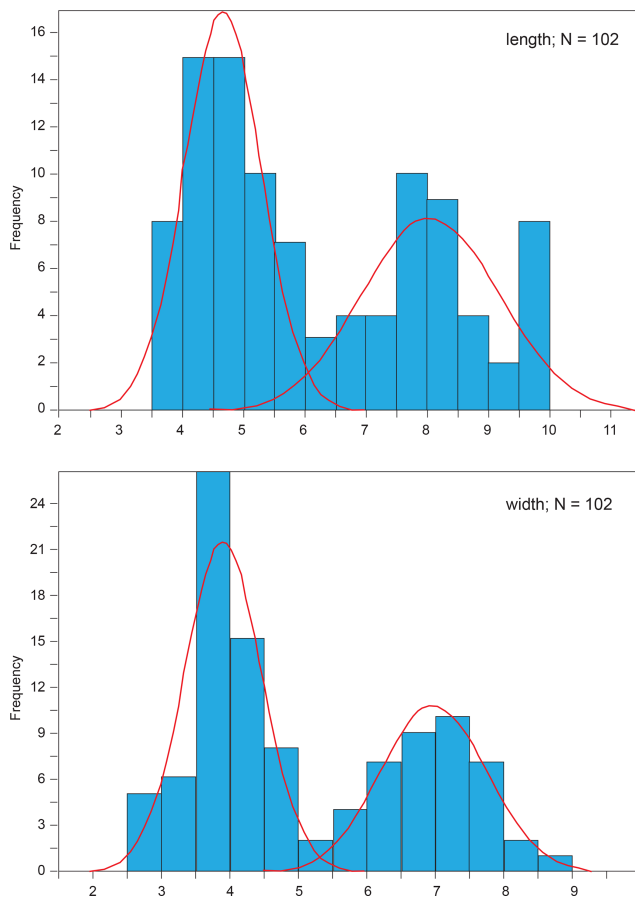


Figure 1. Mixture analysis applied to the length and width of all the analysed coccoliths using PAST 3.01 software (Hammer et al., 2001). The lowest Akaike information criterion (AIC, Akaike, 1973) is obtained with two populations with modes at 4.0–5.0 and 7.5–8.5 μm for coccolith length and at 3.5–4.5 and 6.5–7.5 μm for coccolith width. The number of size classes or bins was chosen in order to obtain each bin at 0.5 μm .

3.2 Coccolith and central-area dimensions

The frequency distribution of coccolith length and width (Fig. 2a and c) for the three analysed species shows that the large morphotype of *Similiscutum*, namely *S. giganteum* sp. nov., can be easily differentiated from *S. finchii* and *S. novum* on the basis of the analysed parameters. However, the boxplot displays important overlapping between *S. giganteum* and *S. finchii* and between *S. finchii* and *S. novum* when taking into account the central-area length and width, which shows a continuum in the size variation (Fig. 2b and d). Based on the Bonferroni–Dunn test, significant differences in the mean sizes between the considered species, both for coccolith and central-area measurements, can be assessed (Table S1 in the Supplement).

The bivariate plot of coccolith length and width and of central-area length and width (Fig. 3a and c) shows that there is a strong covariance between the observed parameters. The mean values with the 10th–90th percentiles of frequency distribution for coccolith length and width for the three analysed species (Fig. 3b) allow an easy differentiation of *S. giganteum* sp. nov. from the cluster composed of *S. finchii* and *S. novum*. However, the bivariate plots of the means for central-area length and width show a continuum in size distribution passing from *S. novum* to *S. giganteum* (Fig. 3d).

3.3 Coccolith geometry

Simple parameters for coccolith geometry, such as the relative proportion of the central area with respect to the coccolith size, are evaluated as the ratios “central-area length over coccolith length” and “central-area width over coccolith width” (Fig. 2e and f; Table S2). The frequency distribution of these two ratios shows a strong overlap for the three analysed species, but the dimensions of the central area relative to the coccolith dimensions of *S. giganteum* sp. nov. are comprised in the lower part of the represented values. There are significant differences in the mean sizes between the considered species for the two ratios, except for *S. giganteum* compared to *S. novum* ($p > 0.0167$), which both possess a relatively smaller central area than *S. finchii* (Table S1).

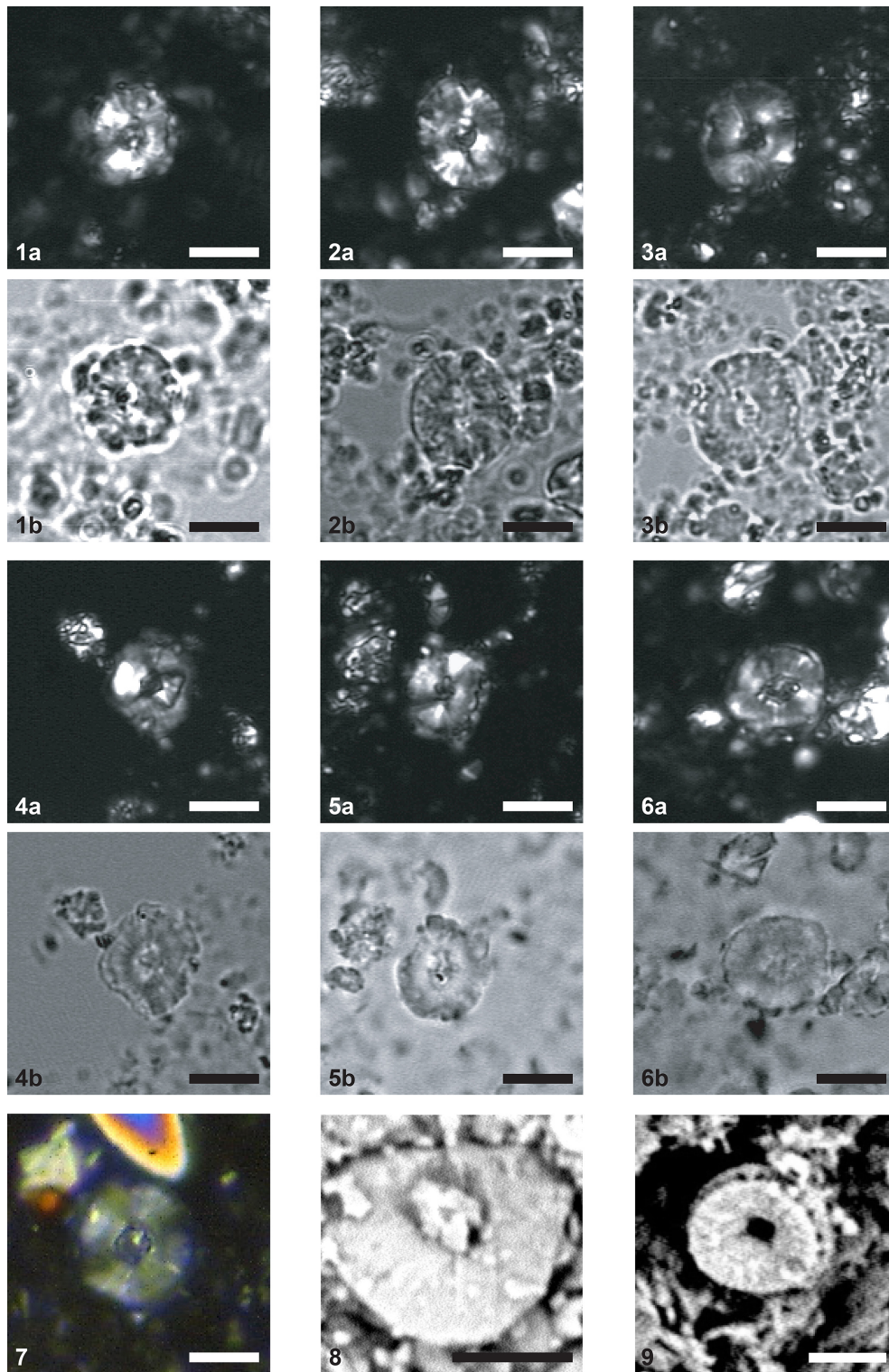


Plate 1. Scale bar = 5 μ m. CPs indicates crossed polars; PPs indicates parallel polars; SEM indicates scanning electron microscope. (1–6) *Similiscutum giganteum* sp. nov. (1a) CPs and (1b) PPs, sample PEN08-Do2003 (Emaciatum Zone), holotype. (2a) CPs and (2b) PPs, sample PEN08-Do2003 (Emaciatum Zone). (3a) CPs and (3b) PPs, sample PEN08-Do2003 (Emaciatum Zone). (4a) CPs and (4b) PPs, sample PEN25bis-To2003 (Polymorphum Zone). (5a) CPs and (5b) PPs, sample PEN25bis-To2003 (Polymorphum Zone). (6a) CPs and (6b) PPs, sample PEN25bis-To2003 (Polymorphum Zone). (7) CPs, sample EMII-7 (Toarcian). (8) SEM, La Cerradura section (Spain), sample CE 34 (Polymorphum Zone). (9) SEM, Borehole FR-210-078, Lorraine Sub-Basin, S Luxembourg, sample K004542 (Falciferum Zone).

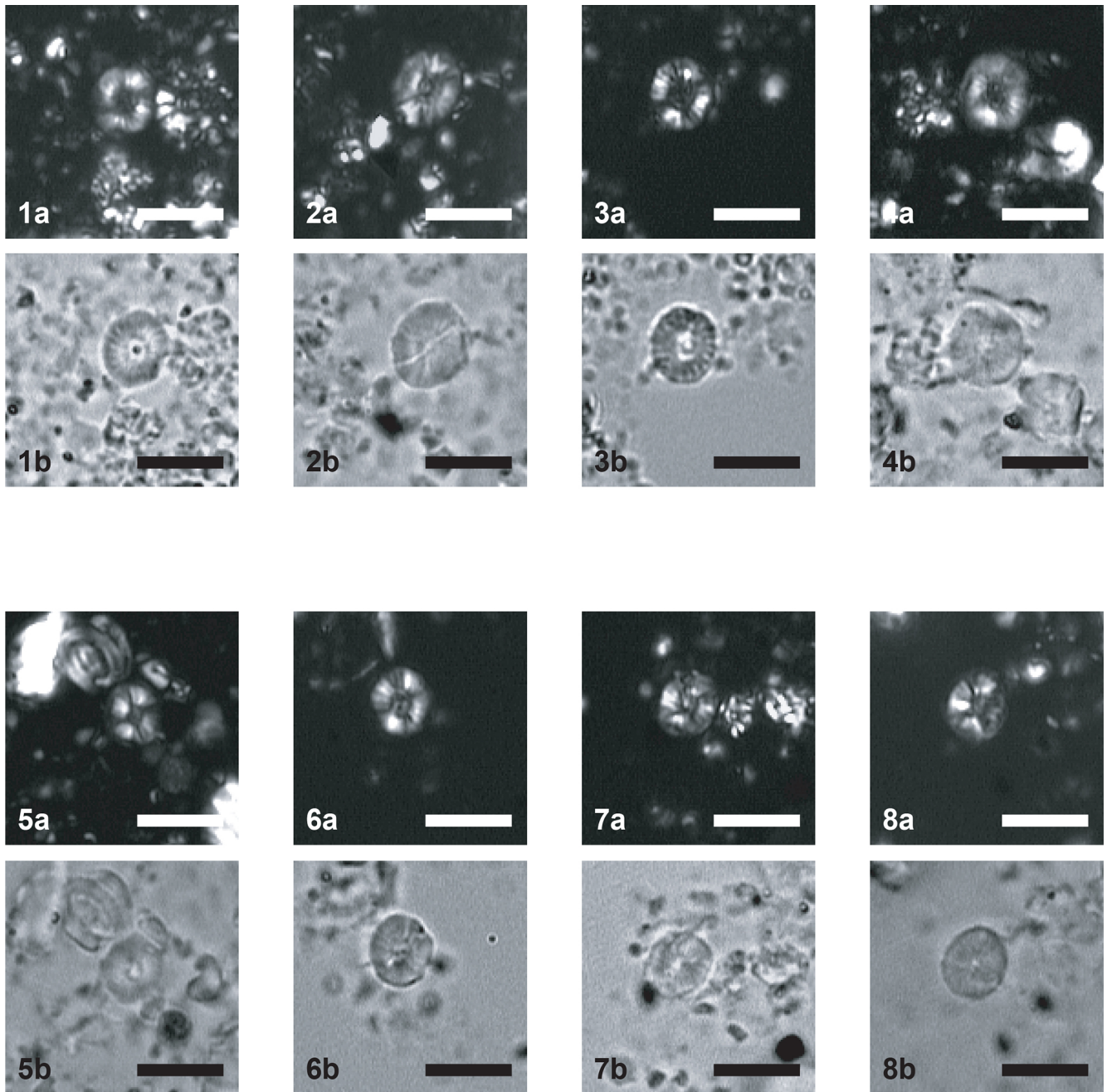


Plate 2. Scale bar = 5 μ m. CPs indicates crossed polars; PPs indicates parallel polars. (1–4) *Similiscutum finchii*. (1a) CPs and (1b) PPs, sample PEN08-Do2003 (Emaciatum Zone). (2a) CPs and (2b) PPs, sample PEN08-Do2003 (Emaciatum Zone). (3a) CPs and (3b) PPs, sample PEN08-Do2003 (Emaciatum Zone). (4a) CPs and (4b) PPs, sample PEN25bis-To2003 (Polymorphum Zone). (5–8) *Similiscutum novum*. (5a) CPs and (5b) PPs, sample PEN25bis-To2003 (Polymorphum Zone). (6a) CPs and (6b) PPs, sample PEN25bis-To2003 (Polymorphum Zone). (7a) CPs and (7b) PPs, sample PEN25bis-To2003 (Polymorphum Zone). (8a) CPs and (8b) PPs, sample PEN25bis-To2003 (Polymorphum Zone).

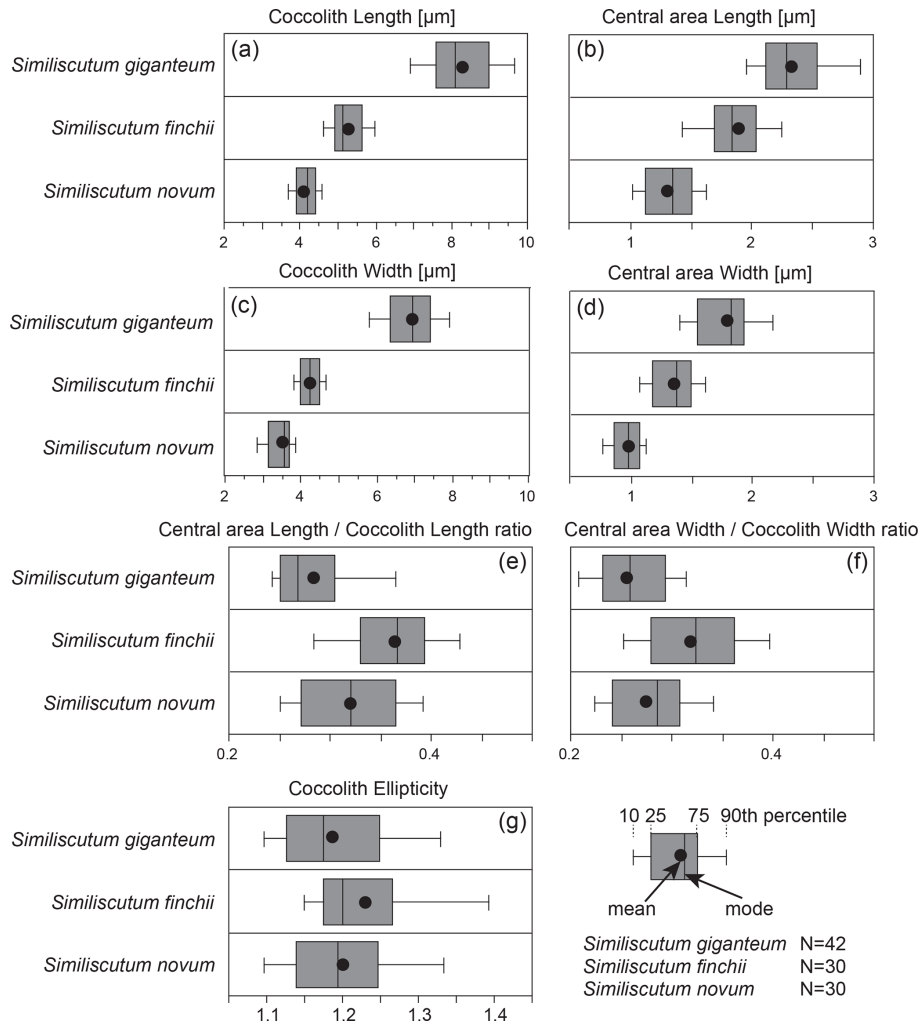


Figure 2. Frequency distribution of coccolith length (a) and width (c) and of central-area length (b) and width (d) of the ratios “central-area length over coccolith length” (e) and “central-area width over coccolith width” (f), along with coccolith ellipticity (g) for the three analysed species. N is the number of measured specimens.

The bivariate plots of the three analysed species (Fig. 3e and f) indicate a fairly good covariance between the ratios “central-area length over coccolith length” and “central-area width over coccolith width”, but the mean central-area dimensions with respect to the coccolith dimensions display a strong overlap between the three species of *Similiscutum*.

Also, the coccolith ellipticity was calculated as the ratio “coccolith length over coccolith width” (Fig. 2g). The frequency distribution of coccolith ellipticity is quite similar for the three analysed species, and no significant differences in the mean ellipticity occur ($p > 0.0167$; Table S1). The bivariate plot of coccolith width versus coccolith ellipticity for the three studied species shows an increase with decreasing coccolith width (Fig. 3g).

3.4 Comparison with previous Biscutaceae measurements

The comparison of the observed size distributions measured in this paper to those published by Mattioli et al. (2004) shows a small overlap in coccolith length and width between *Similiscutum giganteum* sp. nov. and the slightly smaller *Biscutum grande* (Fig. 4a and b). The optical behaviour of these two species is, however, very different. *Similiscutum giganteum* displays a light grey, striated distal shield in optical-microscope crossed polars, without a prominent inner bright “collar”, similar to *S. finchii* and *S. novum*; on the other hand, the *Biscutum grande* uni-cyclic distal shield appears homogeneously dark grey and possesses a white, birefringent “collar” surrounding the central area (Bown, 1987a; Mattioli et al., 2004).

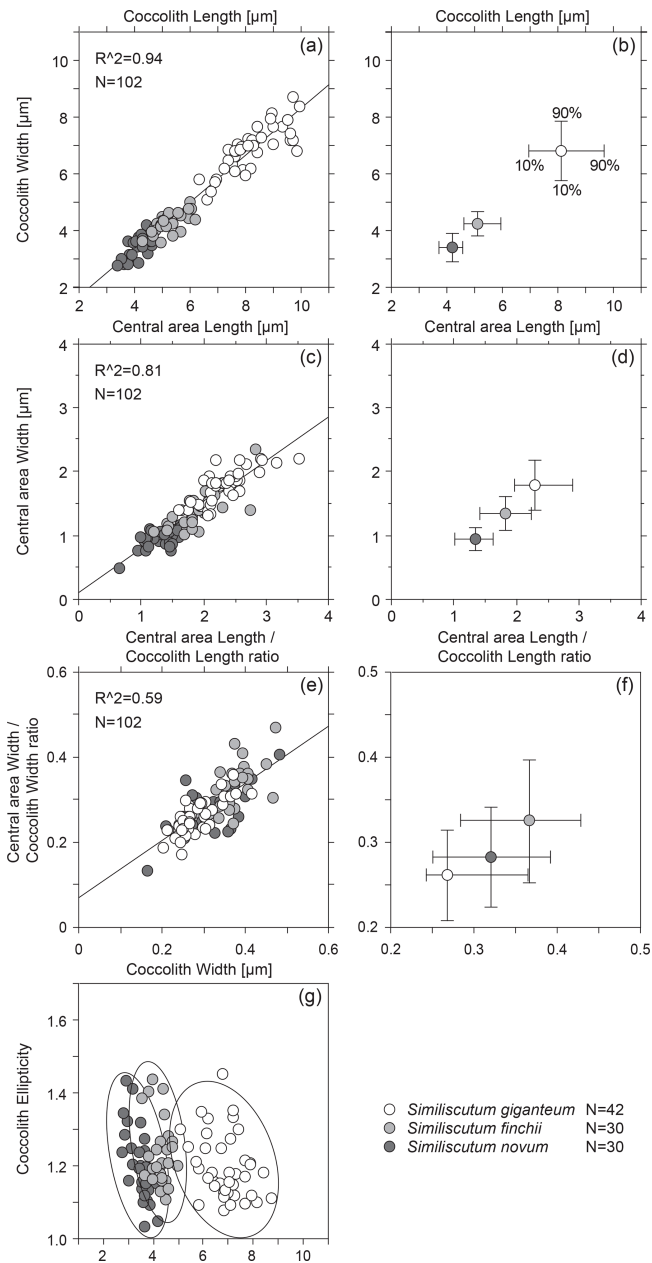


Figure 3. Bivariate plot of coccolith length and width (a) and of central-area length and width (c). The calculated R^2 is quite high. Mean values with the 10th–90th percentiles of frequency distribution for coccolith length and width (b) and for central-area length and width (d) for the three analysed species. Bivariate plot of the ratios “central-area length over coccolith length” and “central-area width over coccolith width” (e) and mean values with the 10th–90th percentiles of frequency distribution for these two ratios (f) for the three analysed species. Coccolith ellipticity versus coccolith width (g). N is the number of measured specimens.

4 Interpretation and taxonomy

New biometric analyses performed on *S. finchii* and *S. novum* and on a large new species of *Similiscutum*, including coccolith length and width and their average values, allow an easy differentiation of the new species from the cluster composed of *S. finchii* and *S. novum*. From a morphological and biometrical point of view, *S. finchii* and *S. novum* show similar features, both being medium-sized, normal to broadly elliptical coccoliths (Mattioli et al., 2004). *Similiscutum novum* is, however, slightly smaller in size and less elliptical and has a smaller central area. In SEM (scanning electron microscope) pictures (like for the holotype; Fig. 5), the central area of *S. finchii* is lenticular, elongated and narrowly elliptical, while in *S. novum* the central area is rather sub-circular (see the holotype in Fig. 5). In LM (light microscope) crossed polars, the central area of *S. finchii* clearly appears elongated and sub-rectangular (Plate 2, figs. 1–4), while in *S. novum* it is rather squared (Plate 2, figs. 5–8). The different stratigraphical ranges reported for *S. finchii* and *S. novum* (Bown and Cooper, 1998; Mattioli and Erba, 1999; Ferreira et al., 2019) are a further argument to keep these two species separated.

The central-area length and width and their means allow the differentiation of the large morphotype of *Similiscutum* from *S. novum* but not from *S. finchii*. The relative proportion of the central area with respect to the coccolith size (ratios “central-area length over coccolith length” and “central-area width over coccolith width”) and the coccolith ellipticity (ratio “coccolith length over coccolith width”) cannot be confidently used as diagnostic parameters for differentiating between these three morphotypes of *Similiscutum*.

Taking into account only size differences, the very large morphotype of *Similiscutum* had been previously considered to be a big morphotype of *S. finchii* (a probable ecophenotype), since they exhibit a similar optical behaviour. In fact, in the literature, a differentiation was made between small and large specimens of *S. finchii* by several authors (e.g. Bown and Cooper, 1998; Menini et al., 2019; Chaumeil-Rodríguez et al., 2022). However, the original diagnosis of *S. finchii* (“A species of *Biscutum* with a large central area and no spine” (Crux, 1984, p. 168)) does not satisfactorily apply to the new species described here, which has a central area smaller than that of *S. finchii* and a cross structure inside it. On the basis of a combination of differences of size and of central-area shape and structure, *Similiscutum giganteum* sp. nov. is introduced here. This is also considerably different in size and geometry from other species of Biscutaceae (for a synthesis, see Fig. 5).

Taxonomic description

Similiscutum giganteum sp. nov.

Plate 1, figs. 1–9

1969 *Palaeopontosphaera crucifera* Prins, pl. 2, fig. 10 (nomen nudum).

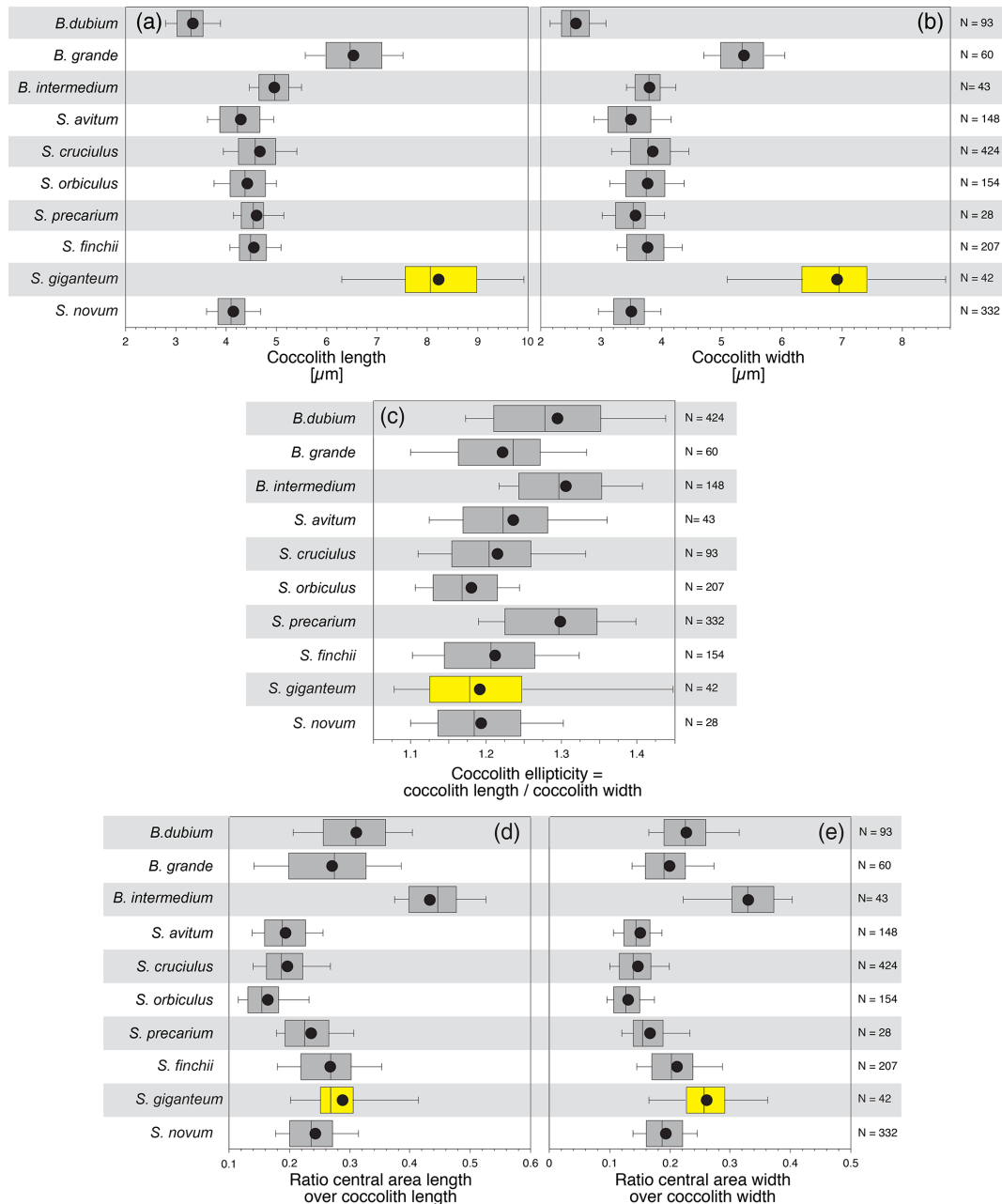


Figure 4. The size and geometry of *Similiscutum giganteum* sp. nov. are compared to similar parameters measured for other Lower Jurassic Biscutaceae after the dataset of Mattioli et al. (2004).

1969 *Palaeopontosphaera veterna* Prins, pl. 2, fig. 9 (nomen nudum).

1987 *Biscutum finchii* Crux, 1984, emend. Bown, 1987a. Bown (partim), pl. 13, figs. 21–22.

1998 *Biscutum finchii* (Crux, 1984, emend. Bown, 1987a) de Kaenel and Bergen, 1993. Bown and Cooper, pl. 4.12, figs. 13–14 (large morphotype).

2002 *Biscutum finchii* (Crux, 1984, emend. Bown, 1987a) de Kaenel and Bergen, 1993. Perilli and Comas-Rengifo, pl. 1, fig. 12.

2006 *Similiscutum giganteum* Mailliot, p. 234, pl. 1, figs. 1–6, unpublished thesis.

2006 *Biscutum finchii* (Crux, 1984) Bown, 1987a. Perilli and Duarte, pl. 1, figs. 7–8.

2008 *Biscutum finchii* (Crux, 1984) Bown, 1987a. Fraguas et al., pl. 1, fig. 10.

2014 *Similiscutum giganteum* Mailliot, 2006. Reolid et al., fig. 6.

2016 *Similiscutum giganteum* Mailliot, 2006. da Rocha et al., fig. 7.9–10.

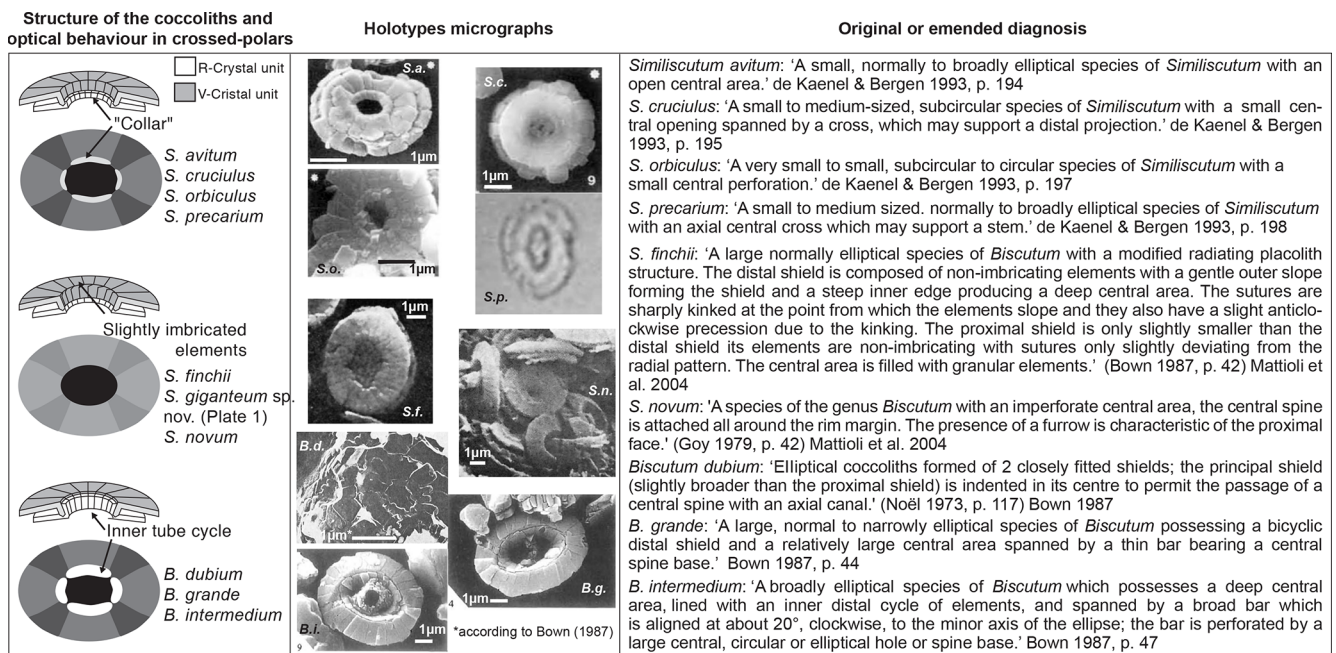


Figure 5. In order to effectively compare *Similiscutum giganteum* sp. nov. to other Lower Jurassic Biscutaceae, the simplified structure in optical microscopes and SEMs (left column), the holotype micrographs (middle column), and the original or emended diagnoses (right column) are reported. An explanation of abbreviations is as follows: *S.a.* = *Similiscutum avitum*; *S.c.* = *Similiscutum cruciulus*; *S.o.* = *Similiscutum orbiculus*; *S.p.* = *Similiscutum precarium*; *S.f.* = *Similiscutum finchii*; *S.n.* = *Similiscutum novum*; *B.d.* = *Biscutum dubium*; *B.g.* = *Biscutum grande*; *B.i.* = *Biscutum intermedium*.

2019 *Similiscutum* aff. *S. finchii* (Crux, 1984, emend. Bown, 1987a) de Kaenel and Bergen, 1993. Menini et al., pl. 2 (large, SN3.57B)

2019 *Similiscutum finchii* (Crux, 1984, emend. Bown, 1987a) de Kaenel and Bergen, 1993. Ferreira et al., pl. 2 (Peniche 61).

2022 Large *Similiscutum* aff. *finchii* (Crux, 1984, emend. Bown, 1987a) de Kaenel and Bergen, 1993. Chaumeil Rodríguez et al., pl. 2, fig. 15.

Derivation of name. From Latin *giganteus*, gigantic.

Diagnosis. A very large, broadly elliptical to elliptical species of *Similiscutum* with a bulky cross spanning the central area. The central area generally exhibits a lenticular shape and is quite reduced compared to the coccolith size.

Holotype. Plate 1, fig. 1 (fig. 1a – CPs: crossed polars, fig. 1b – PPs: parallel polars).

Type locality. Peniche (Ponta do Trovão section), Portugal.

Type level. Sample Pen08-Do2003 (= base of the “transition beds” or “couches de passage” = Bed 15a of Mouterde, 1955) in the uppermost Pliensbachian (Emaciatum ammonite Zone).

Repository: Collections de Géologie de Lyon with the FSL no. 766528.

Description. This coccolith is characterised by a modified radiating placolith structure, similar to *S. finchii* and *S. novum*. The sutures between elements of distal shield often show important kinking. A robust cross spans the lozenge-

like central area. The lenticular central-area shape is particularly visible in proximal view.

Dimensions. Minimum and maximum measured lengths are 6.32–9.93 µm (mean is 8.23 µm); width is 5.08–8.73 µm (mean is 6.91 µm); central-area length is 1.61–3.53 µm (mean is 2.35 µm); central-area width is 1.31–2.21 µm (mean is 1.78 µm).

Holotype (Plate 1, fig. 1) measurements are 8.10 µm (length) – 7.01 µm (width).

Range. Late Pliensbachian (Margaritatus ammonite Zone) to late Toarcian (Meneghinii ammonite Zone) (Ferreira et al., 2019).

Occurrence. Portugal (Peniche, this work); Spain (Camino in Perilli and Comas-Rengifo (2002); La Cerradura in Reolid et al. (2014)); France (Saint-Paul-des-Fonts and Tournadous in Mailliot (2006); ANDRA HTM-102 borehole in Lorraine from personal observation by Emanuela Mattioli (2018); Anse Saint Nicolas in Menini et al. (2019)); Argentina (Picun Leufu in Bown (1987a), Bown and Cooper (1998) and Chaumeil-Rodríguez et al. (2022)).

Discussion. The name *Similiscutum giganteum* was informally used in Mailliot (2006; PhD thesis) but was never formally published. The same name was informally used in Reolid et al. (2014) and da Rocha et al. (2016).

According to biometric data presented in this paper, *Similiscutum giganteum* sp. nov. is one of the largest Biscutaceae of the Lower Jurassic, along with *Biscutum grande* – how-

ever, the latter possesses a peculiarly different extinction pattern in optical microscope and a different structure as far as the inner “collar” is concerned. Bown (1987a) and then Bown and Cooper (1998) have differentiated between small and large morphotypes of *Similiscutum finchii*, with coccolith length ranging from 5.8 to 8.5 µm and coccolith width ranging from 4.8 to 7.0 µm. The large specimens described by these authors significantly differ from the original diagnosis of *Similiscutum finchii* because of their reduced central-area dimensions and more broad elliptical outline of the coccolith (pl. 4.12, figs. 13–14). De Kaenel and Bergen (1993) pointed out an important difference in size between specimens of *Similiscutum finchii* from Portugal, ranging from 5.5 to 8.0 µm, and specimens from Morocco, as small as 5.0 µm. Although not shown, it is not excluded that some of the larger specimens from Portugal noticed by De Kaenel and Bergen (1993) corresponded to *Similiscutum giganteum* sp. nov. Mattioli et al. (2004; p. 16) stated that the size range of *Similiscutum finchii*, as evidenced by biometrics, “... falls at the small end of the range of sizes reported in the literature...”, with a majority of specimens ranging between 4.0 and 5.1 µm. The specimens of *Similiscutum finchii* coming from Portugal and measured in the present work are indeed larger than in other papers, ranging from 4.28 to 6.17 µm, with a mean at 5.23 µm. The presence of over-calcified, robust coccoliths seems to be a common pattern in Peniche (da Rocha et al., 2016). However, the large *S. finchii* mentioned by Bown (1987a), Bown and Cooper (1998) and de Kaenel and Bergen (1993) rather corresponds to *Similiscutum giganteum* sp. nov.

Similiscutum giganteum sp. nov. is also easily distinguished from other *Similiscutum* because of its peculiar, lozenge-like central-area shape, visible in both optical-microscope and SEM images (Plate 1, figs. 4–6 and 8–9). *Similiscutum novum* generally exhibits a small, square to sub-rectangular central-area shape in optical microscopes (Plate 2, figs. 5–8). *Similiscutum finchii* usually displays a sub-rectangular to rectangular, elongated central-area shape (Plate 2, figs. 1–4). *Similiscutum giganteum* sp. nov. also possesses a robust cross in its central area visible in optical microscopes (Plate 1, figs. 1a, 3a, 4b, 5a) and SEMs (Plate 1, figs. 7 and 8). Although this structure can be (partially) broken (e.g. Plate 1, fig. 8), its insertion points on the inner part of the central area are generally well visible. The occurrence of a cross in the central area is a diagnostic parameter for the characterization of this species. The structure of the central area is different for the other *Similiscutum* species. In the description of *Similiscutum novum* (Bown, 1987a; p. 41), it is mentioned that “... the central area of the proximal shield is filled with granular calcite forming a funnel-like structure which protrudes as the spine/tube on the distal side. Only rarely is the central, hollow tube found as a fully developed spine, and this may represent a dimorphic feature or simply be due to preservation...”. The original diagnosis of *Similiscutum finchii* (Crux, 1984; p. 168) reports “... a species of

Biscutum with a large central area and no spine; the central area is filled with irregular granular calcite...”. The emended diagnosis of Bown (1987a; p. 42) states that “... the central area is filled with granular elements...”. De Kaenel and Bergen (1993; p. 877) report for *S. finchii* that “... in cross-polarized light, a bright transverse bar is often observed when specimens are oriented 45° to the polarizing direction; this optical feature results from a thickening of the central plate elements...”. Finally, Mattioli et al. (2004; p. 25) noticed that the central area of *S. finchii* “... can be crossed by simple structures...”. However, the occurrence of a cross in the central area of *S. finchii* is never mentioned in the literature.

Differentiation. *Similiscutum giganteum* can be distinguished from *S. finchii* and *S. novum* by its larger size and by the occurrence of a well-developed cross in the lenticular central area. *Similiscutum giganteum* can be distinguished from the large species *Biscutum grande* by its uni-cyclic distal shield, by its somehow larger size, by the existence of kinking sutures on the distal shield elements and, especially, by the absence of a typical very bright inner tube cycle and of the bridge structure which spans the central area of *B. grande*. The latter may sometimes be characterized by a well-developed cross in its central area, as shown by Menini et al. (2019) (plate 2, LAL-18).

5 Conclusion

Since the 90s, the Jurassic Biscutaceae have been the object of taxonomic revisions, with the introduction of a new genus and several species, which allowed a better biochronological subdivision of the Jurassic period. Since then, the first occurrence of *Similiscutum* has been used to define the lowermost nanofossil zone of the Pliensbachian stage, and this event is now used in the Geologic Time Scale 2020. This paper reports on biometric analyses performed on three species of *Similiscutum*, namely *S. novum*, *S. finchii* and a new, very large morphotype of *Similiscutum* recorded in upper Pliensbachian and Toarcian strata. Because of morphological plasticity shown by *S. finchii*, which presents small and large morphotypes, this new and very large *Similiscutum* was erroneously attributed to *S. finchii*. In fact, as shown by our new biometric data, the original diagnosis of *S. finchii* does not apply to this taxon that has to be considered as a new species. On the basis of combined differences of sizes and of central-area shape and structure, a new taxon, *Similiscutum giganteum* sp. nov., is thus introduced here for designating very large coccoliths, which represent amongst the largest known Jurassic Biscutaceae.

Data availability. All the slides containing nanofossils studied and illustrated in this paper are stored in the Collections de Géologie de Lyon Repository (France) with the label FSL and a number. All the data used in this paper are shown in the Table S1 and S2 in the Supplement.

Supplement. The supplement related to this article is available online at: <https://doi.org/10.5194/jm-42-1-2023-supplement>.

Author contributions. SM, EM and BP performed the sampling. SM performed the sample preparation, light-microscope imaging and biometrics. The species were determined by SM under the supervision of EM. MCR contributed with the taxonomic revision. SM prepared the article and figures with contributions from the co-authors. All authors contributed to writing and editing the article.

Competing interests. At least one of the (co-)authors is a member of the editorial board of *Journal of Micropalaeontology*. The peer-review process was guided by an independent editor, and the authors also have no other competing interests to declare.

Disclaimer. Publisher's note: Copernicus Publications remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Acknowledgements. The authors warmly thank Fabienne Giraud and Serge Elmi for the constructive discussions and review of an earlier version of this paper.

Financial support. Samuel Mailliot was funded by MRT of the French Ministry of Research. Financial support was provided by the Projet ATIP (CNRS PG/MR/No.2002.053).

Review statement. This paper was edited by Kirsty Edgar and reviewed by Angela Fraguas and Jeremy Young.

References

- Akaike, H.: Information theory and an extension of the Maximum Likelihood principle, edited by: Petrov, B. N. and Csaki, F., 2nd International Symposium on Information Theory, Akademia Kiado, Budapest, 267–281, 1973.
- Black, M.: Coccoliths of the Speeton Clay and Sutterby Marl, Proceedings of Yorkshire Geological Society, 38, 381–424, 1971.
- Bown, P. R.: Taxonomy, evolution and biostratigraphy of Late Triassic-Early Jurassic calcareous nannofossils, Spec. Pap. Palaeontol., 38, 1–118, 1987a.
- Bown, P. R.: The structural development of Early Mesozoic coccoliths and its evolutionary and taxonomic significance, Abh. Geol. B.-A., 39, 33–49, 1987b.
- Bown, P. R. and Cooper, M. K. E.: Jurassic, edited by: Bown, P. R., Calcareous Nannofossil Biostratigraphy, British Micropalaeontological Society Publication Series, Kluwer Academic Publishers, 34–85, <https://doi.org/10.1007/978-94-011-4902-0>, 1998.
- Chaumeil Rodríguez, M., Mattioli, E., and Pérez Panera, J. P.: Lower Jurassic calcareous nannofossil taxonomy revisited according to the Neuquén Basin (Argentina) record, J. Micropalaeontol., 41, 75–105, <https://doi.org/10.5194/jm-41-75-2022>, 2022.
- Crux, J. A.: Biostratigraphy of Early Jurassic Calcareous Nannofossils from Southwest Germany, Neues Jahrb. Geol. P.-A., 169, 160–186, 1984.
- da Rocha, R., Mattioli, E., Duarte, L. V., Pittet, B., Elmi, S., Mouterde, R., Cabral, M. C., Comas-Rengifo, M. J., Gómez, J. J., Goy, A., Hesselbo, S. P., Jenkyns, H. C., Littler, K., Mailliot, S., Veiga de Oliveira, L. C., Osete, M. L., Perilli, N., Pinto, S., Ruget, C., and Suan, G.: Base of the Toarcian Stage of the Lower Jurassic defined by the Global Boundary Stratotype Section and Point (GSSP) at the Peniche section (Portugal), Episodes, 39, 460–481, 2016.
- de Kaenel, E. and Bergen, J. A.: New Early and Middle Jurassic coccolith taxa and biostratigraphy from the eastern proto-Atlantic (Morocco, Portugal and DSDP Site 547 B), Eclogae Geol. Helv., 86, 861–907, 1993.
- Favre, E., Escarguel, G., Suc, J.-P., Vidal, G., and Thévenod, L.: A contribution to deciphering the meaning of AP/NAP with respect to vegetation cover, Rev. Palaeobot. Palyno., 148, 13–35, 2008.
- Ferreira, J., Mattioli, E., Sucherás-Marx, B., Giraud, F., Duarte, L. V., Pittet, B., Suan, G., Hassler, A., and Spangenberg, J. E.: Western Tethys Early and Middle Jurassic calcareous nannofossil biostratigraphy, Earth Sci. Rev., 197, 102908, <https://doi.org/10.1016/j.earscirev.2019.102908>, 2019.
- Fraguas, A., Comas-Rengifo, M. J., and Perilli, N.: Pliensbachian calcareous nannofossils of the Santotis section (Basque-Cantabrian Basin, N Spain), Atti della Società Toscana di Scienze Naturali, Serie A, 113, 49–56, 2008.
- Fraguas, A., Comas-Rengifo, M. J., and Perilli, N.: Calcareous nannofossil biostratigraphy of the Lower Jurassic in the Cantabrian Range (Northern Spain), Newsl. Stratigr., 48/2, 179–199, <https://doi.org/10.1127/nos/2015/0059>, 2015.
- Fraguas, A., Comas-Rengifo, M. J., Goy, A., and Gómez, J. J.: Upper Sinemurian–Pliensbachian calcareous nannofossil biostratigraphy of the E Rodiles section (Asturias, N Spain): a reference section for the connection between the Boreal and Tethyan Realms, Newsl. Stratigr., 51/2, 227–244, <https://doi.org/10.1127/nos/2017/0401>, 2018.
- Hammer, Ø., Harper, D. A. T., and Ryan, P. D.: PAST: paleontological statistics software package for education and data analysis, Palaeontol. Electron., 4, 1–9, 2001.
- Hesselbo, S. P., Ogg, J. G., and Ruhl, M. (with contributions by Hinnov, L. A. and Huang, C. J.): The Jurassic Period, in: Geologic Time Scale 2020, Vol. 2, edited by: Gradstein, F. M., Ogg, J. G., Schmitz, M., and Ogg, G., 955–1021, <https://doi.org/10.1016/B978-0-12-824360-2.00026-7>, 2020.
- Mailliot, S.: Production carbonatée pélagique par les nannofossiles calcaires au cours de l'événement anoxique du Toarcien inférieur, PhD thesis, Université Claude Bernard Lyon1, 221 pp. + 2 Annexes, 2006.
- Mattioli, E. and Erba, E.: Biostratigraphic synthesis of calcareous nannofossil events in the Tethyan Jurassic, Riv. Ital. Paleontol. S., 105, 343–376, 1999.
- Mattioli, E., Pittet, B., Young, J. R., and Bown, P. R.: Biometric analysis of Pliensbachian-Toarcian (Lower Jurassic) coccoliths of the family Biscutaceae: intra- and inter-specific variability versus palaeoenvironmental influence, Mar. Micropaleontol., 52, 5–27, 2004.

- Menini, A., Mattioli, E., Spangenberg, J. E., Pittet, B., and Suan, G.: New calcareous nannofossil and carbon isotope data for the Pliensbachian/Toarcian boundary (Early Jurassic) in the western Tethys and their paleoenvironmental implications, *Newsl. Stratigr.*, 52, 173–196, <https://doi.org/10.1127/nos/2018/0476>, 2019.
- Mouterde, R.: Le Lias de Peniche, Direcção geral de minas e serviços geológicos, Lisboa, t. XXXVI das Comunicações dos Serviços Geológicos de Portugal, 1–33 + 4 pl., CDU 551.762.1 (469.323.25), 1955.
- Perilli, N. and Comas-Rengifo, M. J.: Calibration of Pliensbachian calcareous nannofossil events in two ammonite-controlled sections from Northern Spain (Basque–Cantabrian area), *Riv. Ital. Paleontol. S.*, 108, 133–152, <https://doi.org/10.13130/2039-4942/5463>, 2002.
- Perilli, N. and Duarte, L. V.: Toarcian nannobiohorizons from Lusitanian Basin (Portugal) and their calibration against ammonite zones, *Riv. Ital. Paleontol. S.*, 112, 417–434, <https://doi.org/10.13130/2039-4942/6350>, 2006.
- Peti, L., Thibault, N., Clémence, M.-E., Korte, C., Dommergues, J.-L., Bougeault, C., Pellenard, P., Jelby, M. E., and Ullmann, C. V.: Sinemurian–Pliensbachian calcareous nannofossil biostratigraphy and organic carbon isotope stratigraphy in the Paris Basin: Calibration to the ammonite biozonation of NW Europe, *Palaeogeogr. Palaeoclimatol.*, 468, 142–161, <https://doi.org/10.1016/j.palaeo.2016.12.004>, 2017.
- Reolid, M., Mattioli, E., Nieto, L. M., and Rodríguez-Tovar, F. J.: The Early Toarcian Oceanic Anoxic Event in the External Subtropical (South Iberian Palaeomargin, Westernmost Tethys): Geochemistry, nannofossils and ichnology, *Palaeogeogr. Palaeoclimatol.*, 411, 79–94, 2014.
- Suchéras-Marx, B., Mattioli, E., Pittet, B., Escarguel, G., and Suan, G.: Astronomically-paced coccolith size variations during the early Pliensbachian (Early Jurassic), *Palaeogeogr. Palaeoclimatol.*, 295, 281–292, 2010.
- Wiggan, N. J., Riding, J. B., Fensome, R. A., and Mattioli, E.: The Bajocian of the Mid-Jurassic: a critical step in the early Mesozoic phytoplankton radiation, *Earth Sci. Rev.*, 180, 126–146, <https://doi.org/10.1016/j.earscirev.2018.03.009>, 2018.
- Young, J. R., Kucera, M., and Chung, H.-W.: Automated biometrics on captured light microscope images of coccoliths of *Emiliania huxleyi*, edited by: Mokuilevsky, A. and Whatley, R., *Microfossils and Oceanic Environments*, University of Aberystwyth Press, Aberystwyth, 261–280, ISBN 0903878747, 1996.
- Young, J. R., Bergen, J. A., Bown, P. R., Burnett, J. A., Fiorentino, A., Jordan, R. W., Kleijne, A., van Niel, B. A., Romein, A. J. T., and von Salis, K.: Guidelines for coccolith and calcareous nannofossil terminology, *Palaeontology*, 40, 875–912, 1997.