Exceptionally well-preserved Cretaceous microfossils reveal new biomineralization styles

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Calcareous microplankton shells form the dominant components of ancient and modern pelagic sea-floor carbonates and are widely used in palaeoenvironmental reconstructions. The efficacy of these applications, however, is dependent upon minimal geochemical alteration during diagenesis, but these modifying processes are poorly understood. Here we report on new biomineralization architectures of previously unsuspected complexity in calcareous cell-wall coverings of extinct dinoflagellates (pithonellids) from a Tanzanian microfossil-lagerstätte. These Cretaceous 'calcispheres' have previously been considered biomineralogically unremarkable but our new observations show that the true nature of these tests has been masked by recrystallization. The pristine Tanzanian fossils are formed from fibre-like crystallites and show archeopyles and exquisitely constructed opercula, demonstrating the dinoflagellate-affinity of pithonellids, which has long been uncertain.

The interwoven fibre-like structures provide strength and flexibility enhancing the protective function of these tests. The low-density wall fabrics may represent specific adaptation for oceanic encystment life-cycles, preventing the cells from rapid sinking.

Calcification is rare in eukaryotes but has arisen separately as a cell-wall covering in the coccolithophore, foraminifera, and dinoflagellate marine protistan plankton groups[1]. These calcareous shells are widely used as environmental proxies and age-dating tools but despite these widespread geoscientific applications, their biomineralization is relatively poorly understood because it is difficult to maintain most living taxa in culture and their fossils are generally modified by diagenetic recrystallization of fine-scale primary structures. Calcareous shells of dinoflagellates are mostly spherical calcite tests (~10-180 μm) formed by the family Thoracosphaeraceae [2] living in shelf and oceanic surface waters. Around 30 living species and 260 fossil species are known, with most extant forms producing calcareous immotile coccoid cells some of which function as reproductive resting cysts [2,3]. The simple fossil forms that have been included in the calcareous dinoflagellates have wall-architectures with randomly-oblique, radial, tangential, and inclined-radial (pithonellid) crystallographic orientations [4]. Specimens of uncertain biological affinity, lacking diagnostic dinoflagellate characters, have been termed calcispheres or calcitarchs [5] and include the extinct Cretaceous pithonellids. Here we show unsuspected ultrastructural complexity and morphological traits in pithonellid fossils, which have been previously unidentified due to the masking effects of diagenesis. These new structures shed light on the biological affinity of pithonellids and the biomineralogical functionality of their calcareous cell-wall coverings.

Results

Pithonellids – the dominant Cretaceous calcispheres. The extinct pithonellid forms dominate late Cretaceous calcisphere assemblages and can occur in rock-forming abundance [6-8]. Prior to this study, their biomineralogically unremarkable architecture (Fig. 1a) and lack of unequivocal morphological features that characterise dinoflagellates (e.g., tabulation and archeopyles) raised doubts concerning their true biological affiliation [2,3,5]. Most of these forms have appeared to be constructed from two single-layered walls of coarse (>1μm), equidimensional calcite crystallites, although specimens showing variable crystallite size and multiple wall-layers have been observed infrequently and considered to represent ecophenotypic or intraspecific variability [9,10]. Our new observations come from material initially targeted for its preservation of unaltered foraminiferal calcite and used to generate high quality geochemical palaeoclimate proxy records for the Palaeogene [11,12]. Geochemical evidence, but also the extraordinary preservation of coccolithophore calcareous microfossils[13], have demonstrated the exceptional nature of the carbonate preservation in the Tanzanian sediments. Our new results come from Turonian (89–93 Ma) [14] sediments that are 30 million years older and extend this microfossil lagerstätte record into the Cretaceous.

Pithonellid ultrastructure. The Tanzanian pithonellid fossils are incomparably better preserved than any previously documented specimens[15] yet they are assignable to known species based on their gross morphology and crystallographic orientation (Fig. 1b, c). Thus, we can show that the pristine pithonellid walls (Fig. 1d) are constructed from minute (~0.1μm wide), rod-like crystallites, often in groups of similar orientation (Fig. 1e-g), which are arranged in patchwork-like, partly interwoven patterns (Fig. 1f, h). These layers form a submicron-scale lamination of

69 the shell-wall (Fig. 1d, h). Sub-angular, circular and slit-like perforations are typical and often
70 outlined by concentrically-arranged crystallites (Figs. 1g; 2).
71
72 **Discussion**
73 These new observations suggest that most, if not all, previously described pithonellid-type
74 microfossils have been strongly modified by recrystallization and bear little resemblance to their
75 primary architectures. Importantly, we have observed large, taxonomically-diagnostic circular to
76 sub-angular openings (Fig. 1g) and corresponding plate-like coverings (Fig. 1f, i, j). These
77 openings, called archeopyles, and their associated covering plates, called opercula, are
78 characteristic of the immotile cells of dinoflagellates and are related to hatching (excystment) of
79 new motile cells. The presence of distinct apical archeopyles is compelling evidence for the
80 dinoflagellate affinity of the extinct Cretaceous microfossils shown here.
81 The dinoflagellate affinity of these fossils is also supported by the observation of interwoven
82 sub-micron scale crystallite wall-fabrics and the resulting laminated wall-architecture, which is
83 comparable to that seen in certain living dinoflagellates, particularly, *Leonella granifera* (Fig.
84 2d-f). The characteristic interlocking of individual crystals that form the rod-like patterns is seen
85 in both *L. granifera* cysts (Fig. 2e) and pithonellids (Fig. 3a). Study of extant calcareous
86 dinoflagellates has shown that biomineralization is initiated as crystal-like particles of calcite
87 within cytoplasmic vacuoles, which then move to the cell periphery and deposit the crystals in a
88 layer or matrix surrounding the cell [16]. The shell of the coccoid cells of *Leonella* has one matrix
89 [16] and shows no layering while the multi-layered construction of pithonellids suggests a complex
90 matrix consisting of multiple organic layers.

The fine-crystalline interwoven and fibre-reinforced, perforate, laminated fabric of pithonellids (Fig. 3b) forms a shell-construction that provides a balance of strength and flexibility, which is crucial for maximum resilience and thus protection of the cell. Similar woven fibre structures are also seen elsewhere in biominerals, such as, sea urchin tooth calcite [17]. In particular, near identical structure is seen in vertebrate dental enamel fabric (Fig. 3c), where it is largely responsible for the resilience of these structures, made of an otherwise inherently weak material [18] yielding potential applications in biomimetics [17]. Furthermore, such low-density biocalcification is likely a specific adaptation for oceanic encystment life cycles, preventing the cysts from sinking below the photic zone to water depths from which they could not recover following excystment.

Methods

Material and field procedures. The studied specimens were recovered from a borehole drilled during the 2007 phase of the Tanzanian Drilling Project (TDP). Drill Site TDP 22 is situated at 10°04'39.4'' S / 39°37'33.5'' E and recovered lower through middle Turonian sediments (upper *Whiteinella archaeocretacea* and *Helvetoglobotruncana helvetica* planktic foraminiferal biozones) [19]. The sediments are unconsolidated siltstones with a carbonate content typically ranging from 10-15 wt. %. Samples were disaggregated in tap water at the drill site and size-fraction-separated over 20, 63, 125 and 250 µm sieves.

Analyses. Calcispheres were picked from the 20-63 µm size-fraction (5-10 mg per sample) with an eyelash attached to a pen. Microscope magnification was ca. 100x. Scanning electron microscopy (SEM) was performed on a Phillips XL-30 ESEM with a LaB6 electron source. The microscopic analysis comprised four steps. First the whole specimen was documented with

reflected light-optical imaging using a Nikon SMZ1500 binocular light microscope and multiple-plane integrated imaging with NIS Elements™ imaging software. Second, transmitted polarized light optical investigation facilitated identification of the crystallographic wall-type using a Leitz Ortholox binocular microscope after transferring specimens onto a glass slide. An eccentric extinction cross (Fig. 1C) indicates an inclined-radial (pithonellid) crystallographic orientation. Third, specimens were mounted on stubs for SEM imaging of the whole cysts and surface morphology and traits. Fourth, the specimens were opened, using a scalpel, to examine internal wall structure and inner surfaces of the cyst.

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Author Contributions

J.E.W. carried out the palaeontological analyses, imaging, wrote the paper and created figures. P.B. was involved in fieldwork, project organization, palaeontological analyses, wrote the paper and discussed the results with J.E.W.

Additional information

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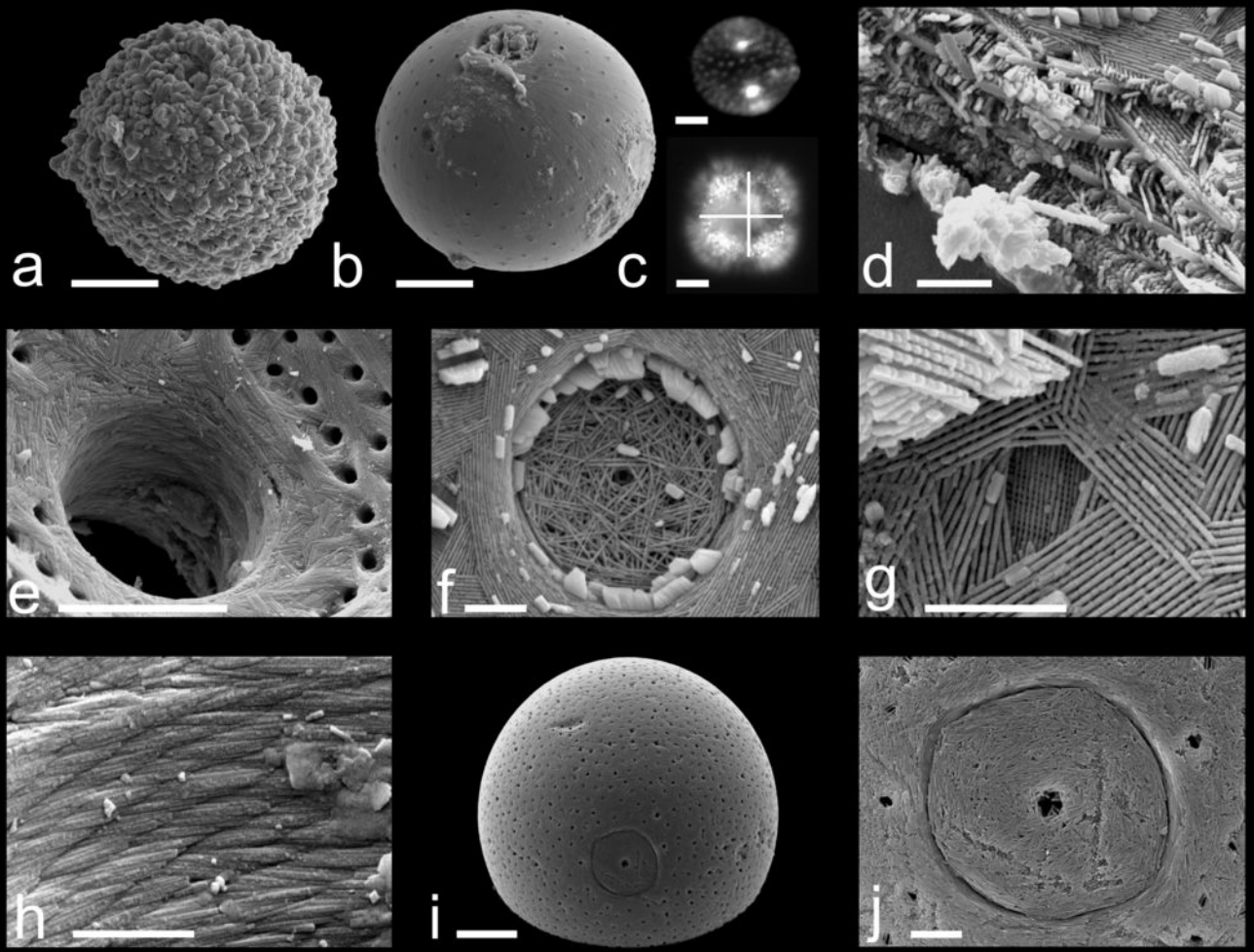
Figure legends

Figure 1 | New biomineralization styles in fossil pithonellids from Tanzanian Drill Site TDP 22. (**a**) *Pithonella* sp.; recrystallized cyst showing spiral rows of prismatic crystals masking primary surface traits. (**b**, **c**) *Pithonella lamellata* (60 specimens investigated); pristine cyst with smooth surface, an archeopyle, spiralling fine striations, and perforations. (**c**) Optical imagery, reflected light (upper) and polarized light (lower). (**d**) Wall-section showing ply-like multiple layers of sub-micron scale, fibre-like crystallites. (**e**) Inner cyst-surface fabric and pore with sieve-like cover. (**f**) Inner cyst-surface with operculum constructed of sub-micron, concentrically interwoven crystallites. (**g**) *Bonetocardiella* sp.; inner cyst-surface with archeopyle, interwoven arrangement of the lamellar wall-crystallite patchwork and circular perforations. (**h**) Archeopyle wall showing rope-like interwoven crystallite bundles and winding architecture. (**i**, **j**) *Pirumella multistrata* (120 specimens investigated), a calcareous dinoflagellate cyst of the randomly-

oblique wall-architecture from TDP Site 22 for comparison; operculum constructed of randomly-oblique, fibre-like crystallites. Scale bars 5 μm.

Figure 2 | Perforation types and comparison with extant calcareous dinoflagellates. (**a**, **b**, **c**) *Bonetocardiella* sp. from TDP Site 22 with sub-angular perforations. (**b**) Wall micro-fabrics in tangential, slightly oblique section. (**c**) Operculum. (**d**, **e**, **f**) *Leonella granifera* (extant species from culture, 8 specimens studied) showing circular perforations. (**e**) Tangential section of wall micro-fabrics consisting of an array of interlocking individual crystals forming a laminated wall, scale bar 2μm. (**f**) Operculum. (**g**, **h**, **i**) *Pithonella ovalis* (238 specimens studied), the type species of the genus *Pithonella*, from TDP Site 22. (**h**) Wall micro-fabrics of parallel fibre-like crystallites and slit-like perforations. (**i**) operculum. Scale bars a-d and f-i 5 μm.

Figure 3 | Ultrastructure of the pithonellid biomineral architecture and comparison to vertebrate enamel. (**a**) Tangential, slightly oblique section of *Bonetocardiella* sp. (TDP Site 22) shows various planes of the wavy lamination (seen in cross-section in figure 1h) of the cyst-wall. Note the interlocking of the individual calcite crystals resembling textile fabrics, scale bar 5 μm. (**b**) Inner shell surface view of the pithonellid fibre-like, and laminated fabric around an enlarged sieve-like perforation formed by orthogonal interlocked crystallites, scale bar 1 μm. (**c**) Cross-ply structure of densely interwoven fibres of rat enamel, scale bar 10 μm (image reproduced courtesy of Dr. Laurie B. Gower, http://gower.mse.ufl.edu/research.html)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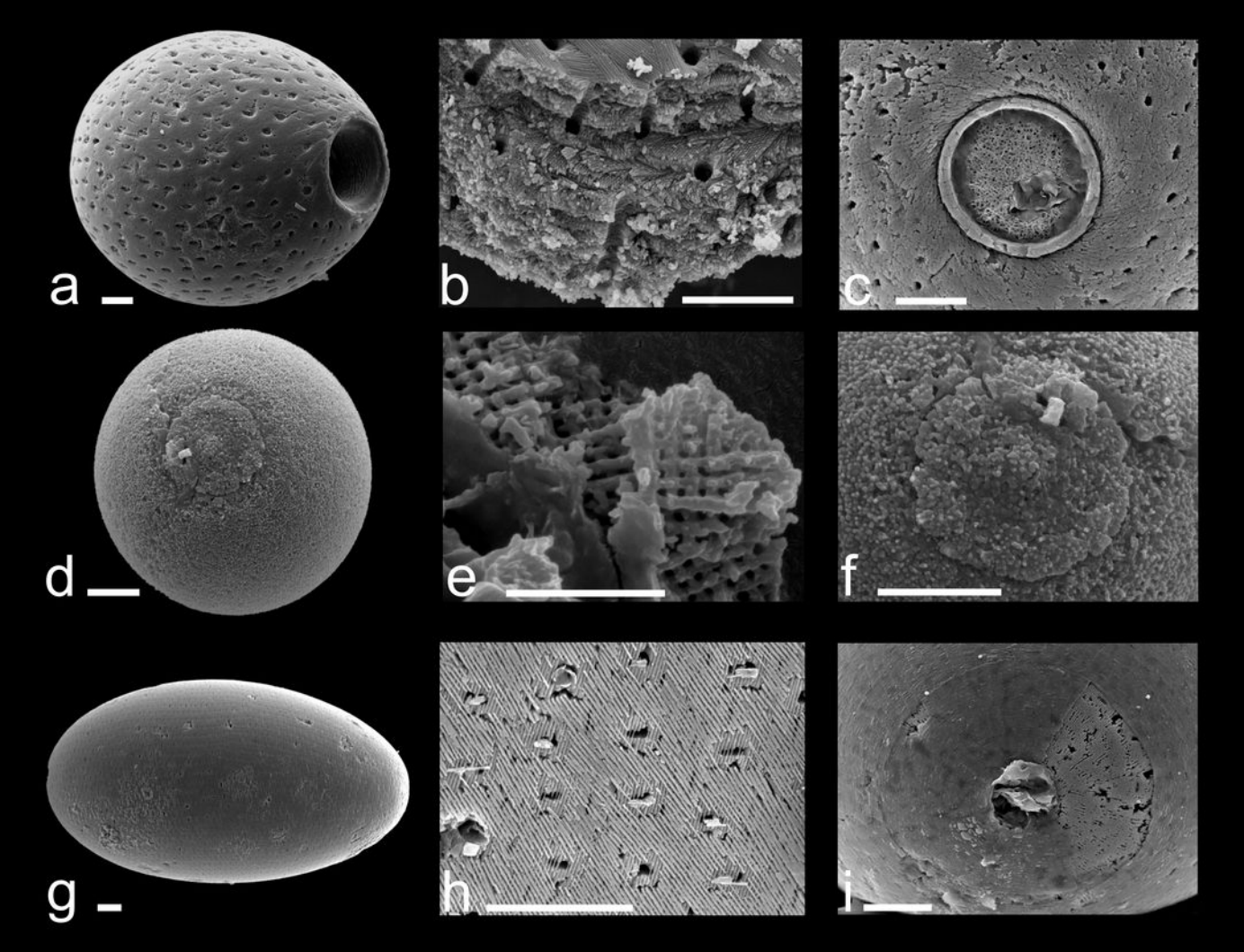
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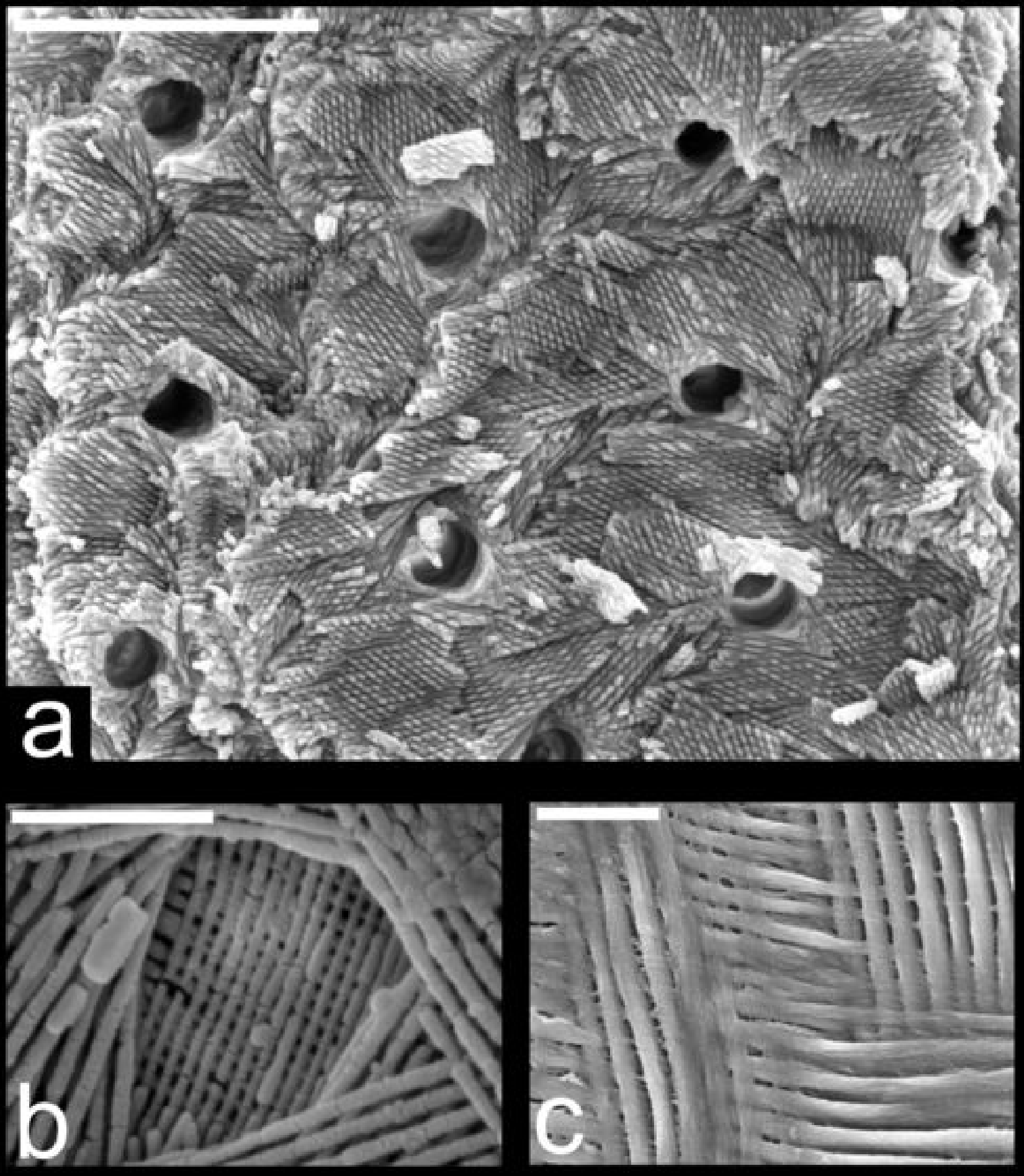
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