https://doi.org/10.5194/egusphere-2025-1840
Preprint. Discussion started: 23 June 2025

The 3D submicron-scale skeletal reconstruction of *Nannoconus* (Cretaceous calcareous nannofossil) - Insights on biomineralization

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Abstract. *Nannoconus* (~5-20 µm) was a major biocarbonate producer in the Early Cretaceous seas (~150-120 Ma). The heavy calcitic skeletons (~200-1400 picogram) of this nannoplankton have contributed massive carbonate accumulations for over ~30 million years. The skeletal microstructure is characterized by an interlocking arrangement of calcitic lamellae spanned around a central canal. The biomineralization process involved in producing the sophisticated skeleton is investigated for the first time. Ptychography X-ray computed tomography (PXCT) with synchrotron radiation is applied to an isolated skeleton, to obtain a 3D set of tomographic images with ~ 40 nm spatial resolution. This 3D set was processed to virtually segment the individual calcitic lamella and reconstruct the full skeleton through constraining different lengths and angles. The lamellae are repetitively stacked in two distinct inclinations, one following the other, and producing segments combined to form the entire skeleton. Individual lamellae were calcified in a "template" of organic layer containing amino acid(s)/biomolecule(s), responsible for creating the interlocking arrangement. Our study of *Nannoconus* provides a simple yet potent approach to the analysis of biomineralized microstructures characterized by the repetitive arrangement of calcitic units as commonly seen in the calcareous nannoplankton.

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1 Introduction

Calcareous nannofossils (~1-30 µm) are biomineralized calcitic remains of marine planktonic unicellular algae (Siesser and Winter 1994) and are abundant in the sedimentary archives. Initially rare, calcareous nannofossils became increasingly abundant in marine sediments to reach an optimum in the Cretaceous, around ~120 Ma (Suchéras-Marx et al., 2019) and they profoundly altered the dynamics of the carbonate and oceanic carbon cycle (Erba, 2006). *Nannoconus* was the major biocarbonate producer in the Early Cretaceous seas (~152-120 Ma) (Bown, 2005) with the largest size (~5-20 µm) and the heaviest mass (~200-1400 picogram) of the calcitic skeleton, contributing to massive carbonate accumulations over ~30 million years. These massively produced biocalcites most likely had significantly modified the marine chemistry upon transfer to the oceanic sedimentary record, with possible consequences on the marine biosphere at a time of important planktonic diversification (Hart et al., 2003; Kooistra et al., 2007). Despite the importance of this group, knowledge of the organism that produced the calcitic skeleton of *Nannoconus* is non-existent. A proper understanding of the biomineralization process from the 3D microstructural arrangement of the skeleton, obtained from a finely-resolved physical characterization, can help to close this gap.

With advances in imaging techniques, several high-resolution characterization methods have been successfully applied to biocalcite produced by extant planktonic algae and preserved as fossils. These techniques include AFM (Henriksen et al., 2004), 3D-FIB-SEM (Hoffmann et al., 2015) and cryo-electron tomography (Walker et al., 2020). In addition, several X-ray scattering methods with synchrotron radiation also have been applied, such as 2D (Suchéras-Marx et al., 2016a) and 3D (Walker et al., 2024) X-ray microfluorescence, X-ray Coherent Diffraction (Beuvier et al., 2019). Ptychographic X-ray computed tomography (PXCT) (Dierolf et al., 2010) is a non-destructive tomographic reconstruction imaging method that can be applied to characterize fossilized biocalcite. In this study, we performed a Ptychography X-ray computed tomography (PXCT) experiment using synchrotron radiation at the SWING Beamline of the SOLEIL (French synchrotron). Our primary focus was on well-preserved skeletons of *Nannoconus* to investigate its microstructural arrangement at the nano-scale resolution. The experiments generated tomographic volumes of the samples with a 3D spatial resolution of approximately 30 nm. Subsequently we analyzed these volumes using image visualization software and successfully reconstructed the complete 3D skeleton of *Nannoconus* for the first time. The knowledge of the 3D microstructural arrangement obtained from the aforementioned reconstruction has been used to provide perspectives on the biomineralization process of the extinct group of calcareous nannoplankton producing such massive skeletons i.e., *Nannoconus*.

2 The *Nannoconus* skeleton

Nannoconus is a Genus (Kamptner, 1931, emend. Farinacci, 1964) belonging to the Family of *Nannoconaceae* (Reinhardt, 1966) (Table C1 in the Appendix). This Family is included in the Order of *Braarudosphaerales* (Aubry, 2013) by Lees and

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Bown (2016). This Order was originally introduced for Cenozoic (~66 Ma to present) coccoliths, another group of calcareous nannofossils.

2.1 Morphology

Different species have been described for the Genus *Nannoconus* and are characterized by different shapes of the skeleton. These include cylindrical, globular, hour-glass and conical (Figs. 1a-1d). Van Niel (1993) introduced several terms to describe it. Some of these terms are used in this study (Figs. 1e and 1f). The *wall* encloses the *central canal* which runs along the longitudinal axis of the skeleton with two terminal openings known as *apertures*. Several morphogroups of *Nannoconus* have been defined based on the general shape and size of the skeleton (Brönnimann, 1955; Bouché, 1965; Derès and Achéritéguy, 1972; Aubry, 1974), the shape of the central canal (Brönnimann, 1955; Bouché, 1965), the thickness and construction of the wall (Brönnimann, 1955; Bouché, 1965; Derès and Achéritéguy, 1972; Aubry, 1974), the shape and the size of the terminal openings (Brönnimann, 1955; Bouché, 1965; Derès and Achéritéguy, 1972; Aubry, 1974).

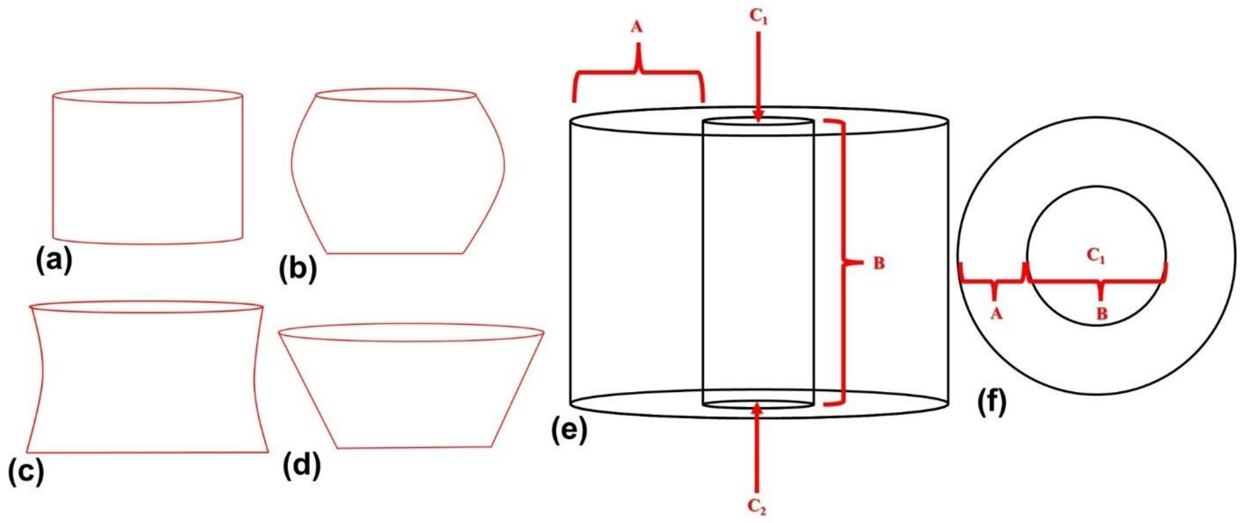

Figure 1: Schematic diagrams illustrating four different skeletal morphologies of four different species of *Nannoconus*. These are (a) cylindrical, (b) globular, (c) hour-glass, and (d) conical as observed in *Nannoconus truittii*, *Nannoconus globulus*, *Nannoconus abundans*, and *Nannoconus steinmannii* respectively. (e and f) Longitudinal and transverse views, respectively, of the schematic illustration of a cylindrical skeleton. In this figure, A and B represent the wall and the central canal respectively. C_1 and C_2 are the two terminal apertures. It should be noted that the shape of the central canal differs with the general morphology of the skeleton.

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2.2 Microstructure

The wall of *Nannoconus* consists of interlocking sub-rhombohedral (Van Niel, 1994) to semi-circular plates (Stradner and Grun, 1973) (length ~0.5-1 µm, thickness ~0.1-0.5 µm) (Kamptner, 1931; Trejo, 1960; Derès and Achéritéguy, 1972; Van Niel, 1993, 1994). Two distinct types of plates (green and yellow line in Fig. 2a) build the skeleton. Each plate is characterized by a specific inclination, measured from a horizontal line (blue line in Fig. 2b). These nannofossils have been defined by Aubry (2013) as "homococcoliths, i.e., coccoliths consisting of identical, imbricated segments that are stacks of lamellae of similar shape". Here, the term "lamellae" refers to the plates of Van Niel (1993) and this term is retained in this work. The term "homococcolith" has been replaced as "micalith" by Aubry (2025).

Based on 2D scanning electron microscopy (SEM) studies, some authors (Aubry, 1974, 2013, 2025; Van Niel, 1993) attempted to explain the 3D microstructural arrangement. Currently two different concepts of skeletal construction are proposed. The first concept is based on a layering of two types of plates (Van Niel, 1993; Lees and Bown, 2016), the second concept is based on an organization of lamellae in segments (Aubry, 2013, 2025) of *Nannoconus*. They can be summarized as the following two models:

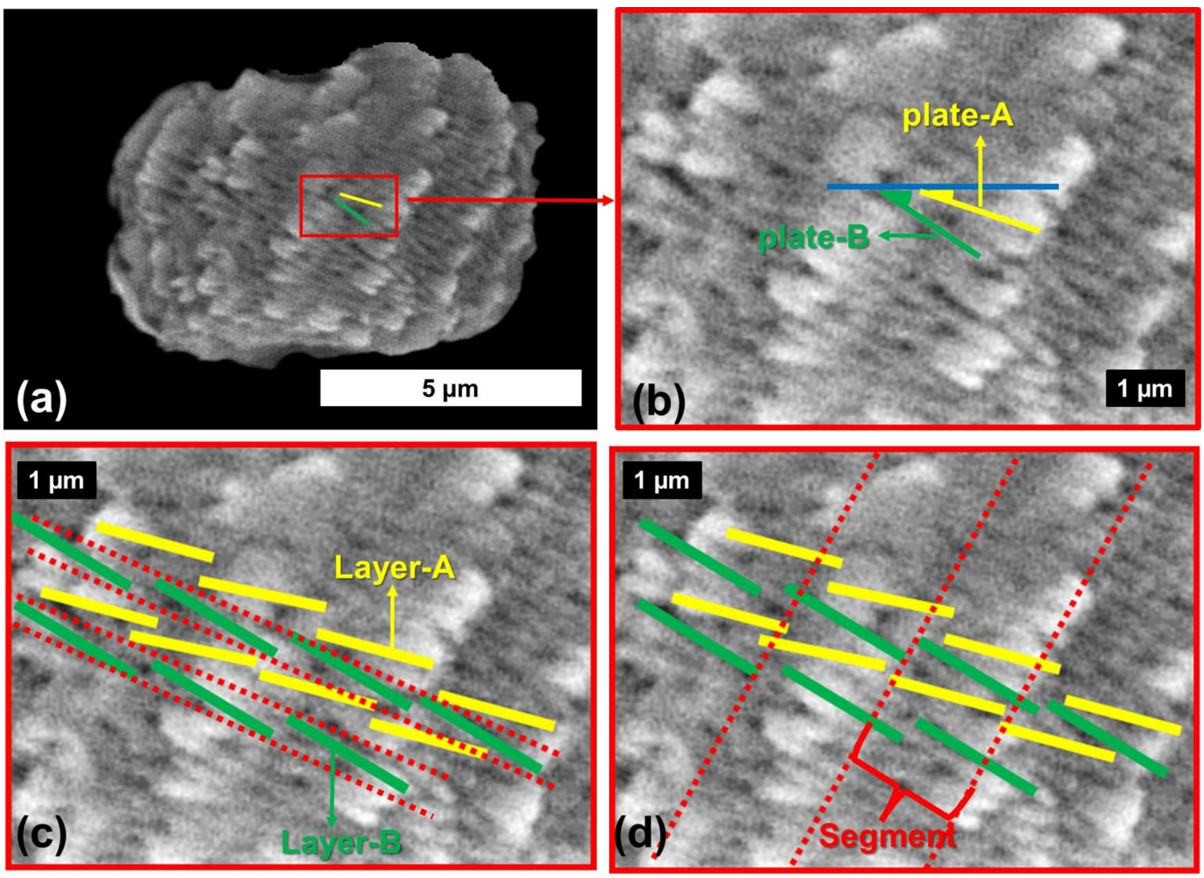

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Figure 2: Scanning electron microscopic (SEM) image of *Nannoconus globulus*. (a) Longitudinal view. The solid green and yellow lines inside the red box highlight the two distinct types of plate-A and -B. (b) Closer view of the plate-A and plate-B shown in (a), with the inclinations measured from the blue horizontal line. (c) Arrangement of plates -A and -B in two layers (-A is yellow and -B is green). The red-dotted line separates the two consecutive layers. (d) Oblique arrangement of the segments composed of alternating plate-A (yellow) and -B (green). Consecutive segments are individualized by red dotted lines. "Plates" are used following the terminology of Stradner and Grun (1973) and Van Niel (1993); they have been referred to as "lamellae" by Aubry (2013). The term "lamellae" is retained throughout this work.

2.2.1 Model-1

In Van Niel (1993), the plates with a lower angle of inclination (yellow) are termed plate-A and the plates with a higher angle of inclination (green) are termed plate-B. Plate-A and -B form two distinct types of layers (Fig. 2c) named as layer-A and layer-B respectively. The skeleton can be understood as a combination of these two types of spiral layers of plates alternatively arranged one after another. This can be explained in Fig. 2c: first, a layer-B (green lines) is placed, followed by a layer-A (yellow lines). Over layer-A, another layer-B is again placed. Delimitation of layers is marked by red dotted lines (Fig. 2c). Van Niel (1993) distinguished nine different morphogroups (Table C1 in the Appendix) of *Nannoconus*, based on the various inclination(s) of plate-B and the width(s) between two consecutive layers of plate-A.

In a SEM image of the species *Nannoconus globulus* (selected for the present study), it is possible to count six plates with two distinct angles of inclinations in the part where the plates are well-preserved (Figs. B1a and B1b in the Appendix). This part constitutes about half of the circular aperture (180°). Thus, the total number of plates in the full circular aperture (360°) would be =12, with six plate-A and six plate-B. Considering that each of these plates belongs to a separate layer, there are six layer-A and six layer-B, i.e., a total of 12 layers in the whole skeleton.

2.2.2 Model-2

As suggested by Aubry (2013, 2025), the skeleton is a combination of identical, imbricated segments. These segments are formed by stacking lamellae (Fig. 2d) of similar shape (Fig. 12d in Aubry, 2025). Lamella-B (green line) and lamella-A (yellow line) are organized two by two to form duos. Duos of lamellae are stacked to compose a segment. Therefore, the spiral combination of several such segments forms the entire skeleton. Three such segments are delineated by red dotted lines in Fig. 2d.

In the SEM image (Fig. B1c in the Appendix) of *N. globulus* described above (in model-1), six segments in the part where the skeleton is well-preserved, can be detected. As this part constitutes about 1/2 of the circular aperture (180°), the total number of segments in the full circular aperture would be 6*2 = 12. Therefore, the total number of the segments in the full skeleton of *Nannoconus* = 12.

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It should be noted that these two concepts only explain the skeletal microstructure and do not provide any information about the parameter(s) controlling the shape of the skeleton. To address the validity of the two models and also to decipher the parameter(s) influencing the skeletal shape and microstructure from the 3D skeletal reconstruction, a synchrotron-based PXCT was conducted.

2.3 Stratigraphy

Nannoconus spans from the base of the Lower Cretaceous through Campanian (Perch Nielsen, 1985). However, it is only abundant until the end of the Aptian which is the end of the climax of *Nannoconus* (Erba, 1994). This represents a duration of ~30 Myr, during which the skeleton exhibits a great morphological diversity (Derès and Achéritéguy, 1980).

3 Materials and Methods

3.1 Material

Six specimens representing five different species of *Nannoconus* have been used for the Ptychographic X-ray Computed Tomography (PXCT) experiment. They are from the Aptian (Covington and Wise, 1987) calcareous sediments of the DSDP Leg-93-Site 603 (Atlantic Ocean, lower continental rise of Cape Hatteras) core 44, interval 115-116 cm. They have been selected for their good state of preservation as determined through scanning electron microscopy (SEM images of *Nannoconus* are shown in Fig. B2 in the Appendix). We focus our study on the species *Nannoconus globulus* (Brönnimann, 1955) to illustrate the microstructural arrangement and the 3D reconstruction of the skeleton. This globular skeleton exhibits a well-defined interlocking pattern of the lamellae.

3.2 Sample Preparation

The PXCT experiment required the picking of a single specimen of *N. globulus* with a silica needle (Figs. B3a and B3b in the Appendix) (Chowdhury et al., in preparation, modified from Suchéras-Marx et al., 2016b). The needle was then placed in a metal holder (Fig. B3c in the Appendix), directly placed at the experimental station of the beamline.

3.3 Ptychography computed X-ray tomography (PXCT)

Ptychography is an advanced X-ray imaging technique that involves capturing a series of far-field diffraction patterns from overlapping areas of a sample. These diffraction patterns are then used in a computational phase retrieval algorithm to reconstruct a high-resolution two-dimensional image. In the field of PXCT, ptychography is combined with tomography. This process involves performing ptychographic experiments at different tomography angles, which produces highly spatially resolved detailed tomographic projections. As a result, PXCT offers a thorough three-dimensional (3D) representation of the microstructure of the sample, achieving nanometric resolution. This enhances the ability to analyze complex materials at a microstructural level. After obtaining the 3D images, it is essential to examine the resulting high-resolution tomographic volume of biocalcite using advanced 3D image analysis techniques. Methods such as image segmentation and visualization (Reznikov et al., 2020) are necessary to extract relevant microstructural properties from the sample.

The experiments were performed at the SWING beamline at SOLEIL-Synchrotron, Paris, France. The photon energy was of 8.00 keV, and the sample-to-detector position was set to 6.484 m. The ptychography experiments were carried out with a beam size of about 4 µm at the sample position and a scanning step size of 1.0 µm. A number of 164 positions were collected and distributed as concentric circles within a rectangular field of view of 20 µm x 10 µm (HxV). We used a region of interest of 1000 x 1000 pixels of an EIGER-4M in-vacuum detector from DECTRIS with a pixel size of 75 µm. For the PXCT experiments (Dierolf et al., 2010), 632 tomographic projections were acquired between 0° and 180°, with exposure time of 100 ms, as 8 interlaced sub-tomograms. The resulting diffraction patterns were transformed into real space projections using the Ptycho-Shelves (Wakonig et al., 2020) suite from the cSAXS team at Paul Scherrer Institute, with a starting probe model previously determined experimentally using a Siemens Star ptychographic scan. The final volume was determined from the tomographic projections using the Matlab tomography package within PtychoShelves.

4 Results

4.1 Descriptive parameters

The combination of the tomographic image slices, obtained from PXCT, reveals the external and internal views of the *N. globulus* skeleton. These observations of the *Nannoconus* are presented here to understand the shape of the skeleton and the arrangement of lamellae and to propose a set of standard parameters that can be used to describe other *Nannoconus* morphogroups.

In the following, cylindrical coordinates are used to describe the skeletal microstructure in terms of the positions and orientations of the constituting lamellae (Figs. 3a-3d). The *z*-axis is parallel to the longitudinal view of the skeleton, passing through the central canal. The Radius, *r*, corresponds to a vector going from the center of mass of the single lamella to the center of the central canal, and it is perpendicular to the *z*-axis. For the purpose of this geometric description, the lowest and the highest points of the skeleton in longitudinal orientation are defined as base and apex, respectively (Fig. 3a). The total vertical length between the base and apex is given by L (Fig. 3a). The length of the radius (r) is measured from the central axis to the end of the wall (Fig. 3a). It changes throughout the whole skeleton from the base to the apex. Starting from the base, the radius increases, reaching a maximum and decreasing towards the apex, giving the skeleton a globular shape. This also indicates that the radius value is maximum at 2/3rd of the length (measured longitudinally) between the base and the

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apex. The minimum value of the radius is observed at the base of the skeleton (Fig. 3a). Two angles define the spatial orientation of each lamella. The first angle was previously defined by Van Niel (1993), based on the SEM images and is Inclination (*δ*). In the cylindrical coordinates described before, it corresponds to the angle (Fig. 3b) formed between the lamella and an axis corresponding to the radius (blue line in Fig. 3b). Two sets of lamellae with different inclinations are shown in Fig. 3c: the lamella with low inclination (δ_A, yellow line given by VW), is named lamella-A. The lamella with high inclination (δ_B, green line given by XY), is termed as lamella-B (Fig. 3c). Lamella-A + lamella-B formed twin lamellae. The second angle (Fig. 3d), called Tilt (*τ*), corresponds to the angle formed between a lamella and an axis of rotation perpendicular to the radius (red line in Fig. 3d). Tilt values are of opposite sign for lamellae placed in the upper (high z) or lower (low z) parts of the skeleton (Fig. 3d). As mentioned above, the arrangement of the lamellae has been described in two models in the literature. Here, we will show that the components (the layers in model-1 and the segments in model-2) of the two models can be verified using the abovementioned radius and angles.

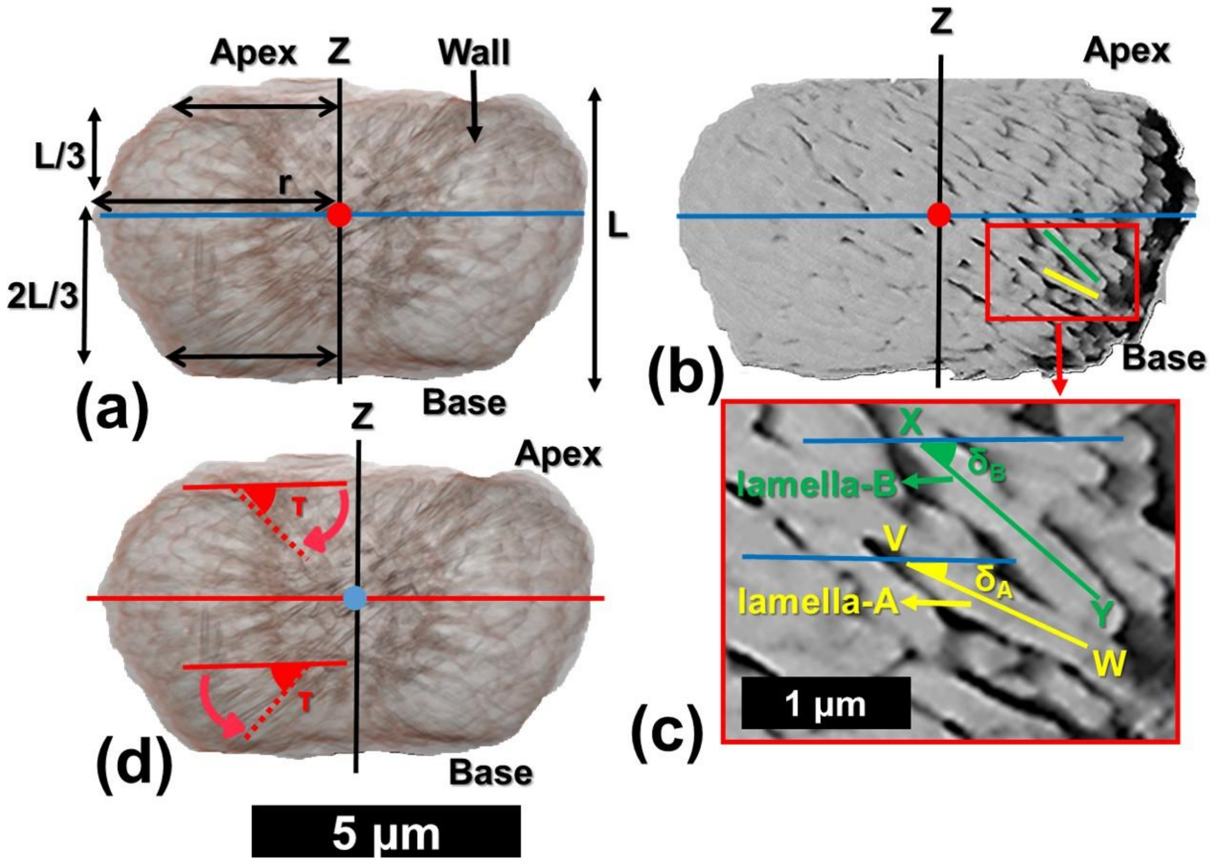

Figure 3: Description of the parameters for the reconstruction of the 3D skeletal microstructure of *N.globulus* using the results of the PXCT experiment. (a) Internal view of *N. globulus.* The microstructure is interpreted using a cylindrical coordinate system.

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The red dot indicates the radial axis (perpendicular to the plane of the paper), and the blue line represents an axis perpendicular to the radius. The z-axis, marked by the black line, passes longitudinally through the central canal. The length of the red arrows corresponds to the radius, measured from the z-axis to the end of the wall of the skeleton. The radius increases initially from the base for 2/3rd of the length (L) of the *Nannoconus*, and then decreases towards the apex for the remaining 1/3rd of the length (L). (b) External view of the same specimen of *N. globulus*, in the cylindrical coordinate system. (c) Magnified skeletal microstructure of the part delineated by the red box in (b), illustrating the two lamellae types: lamella-A (yellow) and lamella-B (green). Their inclinations, δ_A and δ_B respectively, correspond to the angles measured between the blue line (axis perpendicular to the radius) and each lamella. (d) The angle measured between the lamella and the red line (radial axis) defines the tilt. The tilts of the lamellae are opposite in direction; for the lamellae at the base (lower z) and the apex (upper z).

4.2 Segmentation of a lamella

The first step in our analysis of the PXCT data was to isolate one lamella (Figs. 4a-4g). The experiment resulted in a series of tomographic image slices for the whole skeleton. Despite having a satisfactory spatial resolution of ~30-40 nm, the lamellae are sometimes difficult to segment (i.e., virtually separate) from the images. This is caused by internal overlapping of the lamellae and by the limited spatial resolution at the same scale of the lamella thickness. To overcome this problem, all the images were filtered using Contrast Limited Adaptive Histogram Equalization (CLAHE) (Reza, 2004). This increased the contrast of the images and hence helped to see more clearly the boundaries between the lamellae. After this contrast enhancement, a single lamella was segmented by selecting by hand regions of interest for each slice. Four such slices are shown in Figs. 4a-4d. This results in a 3D volume of the lamella that was exported and used to reconstruct the structure of the skeleton using our models (Figs. 4e and 4f). Given the high amount of data, the convex hull (Table C1 in the Appendix; Ky, 1959) of the 3D volume of the segmented lamella (Fig. 4g) was generated using Python routines and utilized for the reconstruction of the models. The segmented lamella is of triangular, flat shape, ~2.00 µm long and ~0.50 µm thick.

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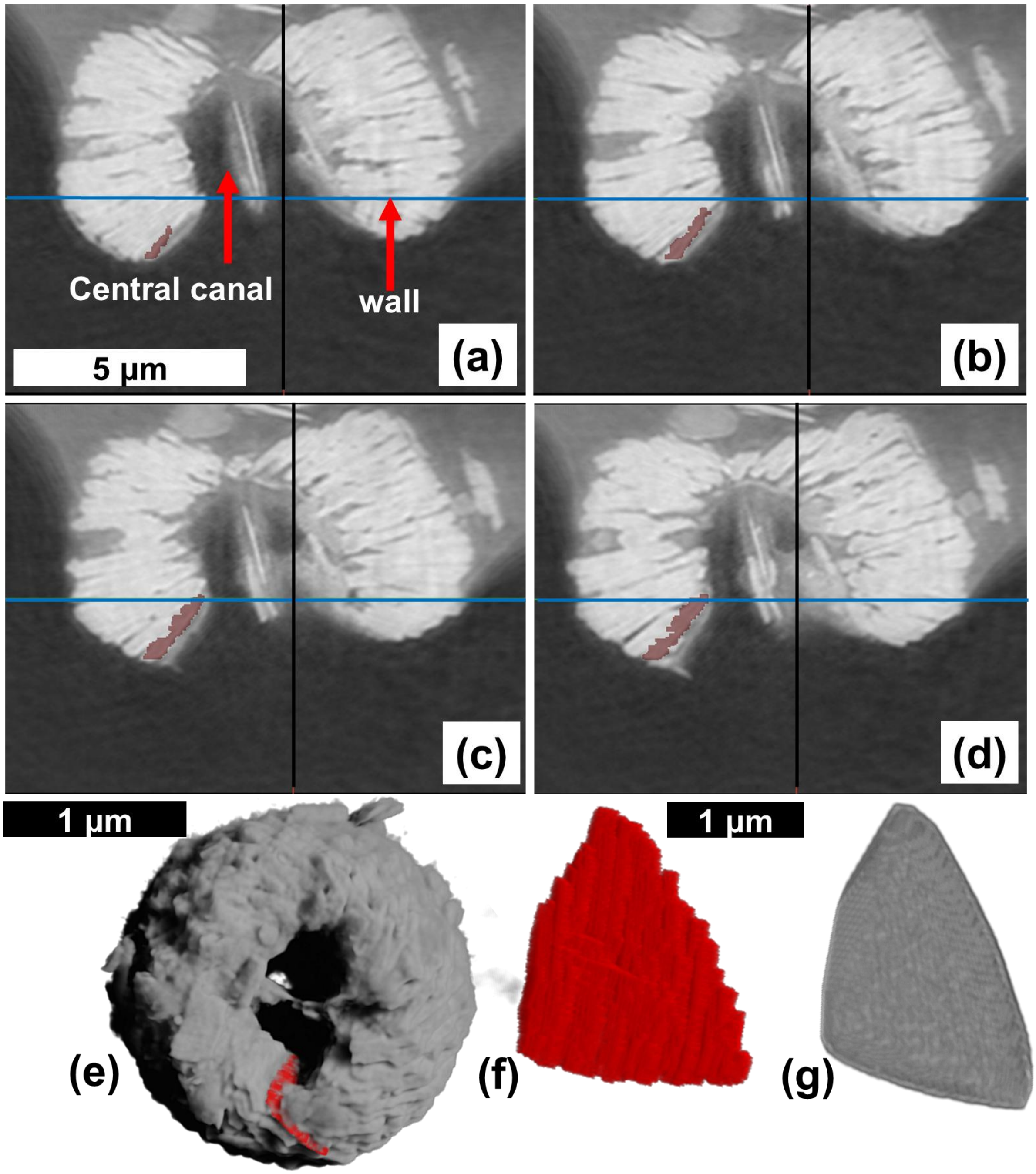

Figure 4: Procedures describing the segmentation/virtual separation of a lamella from the result of the PXCT analysis of *N. globulus*. (a-d) Four consecutive tomographic images obtained from the PXCT analysis of *N. globulus* with the wall and central canal. The part present within the central canal is possibly a lamella detached from the wall. (e) The marked lamella (red colour) is presented within the skeleton. (f) Isolated individual lamella after the segmentation. (g) Convex hull of the lamella, with smooth edges and surfaces. Convex hull (Table C1 in the Appendix) is the mathematical representation of an envelope covering the segmented lamella.

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4.3 Reconstruction

We developed a code using Python script (Van Rossum and Drake, 2009) directly in a console attached to the image visualization software to reconstruct the skeleton using the segmented lamella. The process of the skeletal reconstruction from the segmented lamella can be described as:

4.3.1 Generation of inclination and tilt

The lamella (given by ABC in Fig. 5a), which has a triangular, flat shape, is rotated in two perpendicular directions in a cylindrical coordinate system to set values for inclination and tilt. The procedure is as follows:

1. The rotation with the radial axis as the axis of rotation generates the inclination, δ, of the lamella (Figs. 5a and 5b).

2. The rotation with the axis perpendicular to the radial axis (Figs. 5a and 5c) as the axis of rotation generates the tilt, τ, of the lamella.

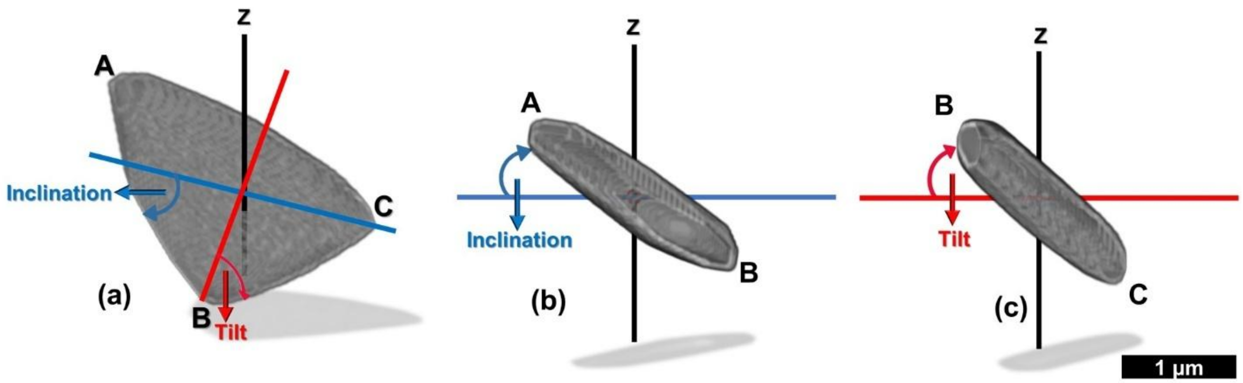

Figure 5: Generation of the inclination and tilt in the segmented lamella. (a) The combination of two perpendicular rotations in the cylindrical coordinate, creating the inclination and tilt in the segmented lamella. (b) The rotation of the lamella, where the radius (red line) is the axis of rotation, creates inclination, and the rotation angle is measured between the lamella and the blue line. (c) The rotation of the lamella, where the axis perpendicular to the radius (blue line) is the axis of rotation, creates tilt, and the rotation angle is measured between the lamella and the red line. It should be noted that the axes of rotation for creating inclination and tilt are perpendicular to the lines from which the individual angle of inclination and tilt are measured.

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4.3.2 Formation of points in spiral axes

The skeleton being conceptualized as a combination of spiral layers/segments, it is now necessary to create the layers/segments utilizing the segmented lamellae. A series of points forming spiral axes was generated to place individual lamellae, creating layers/segments. The points of a series are defined following cylindrical coordinates (Fig. 6a):

i) The radius, r, given by the distance from the central axis to the center of mass of the lamella.

ii) The azimuth, ɵ, given by the angle around the central axis.

iii) The vertical distance, z, perpendicular to the radius and along the axis of rotation of the azimuth.

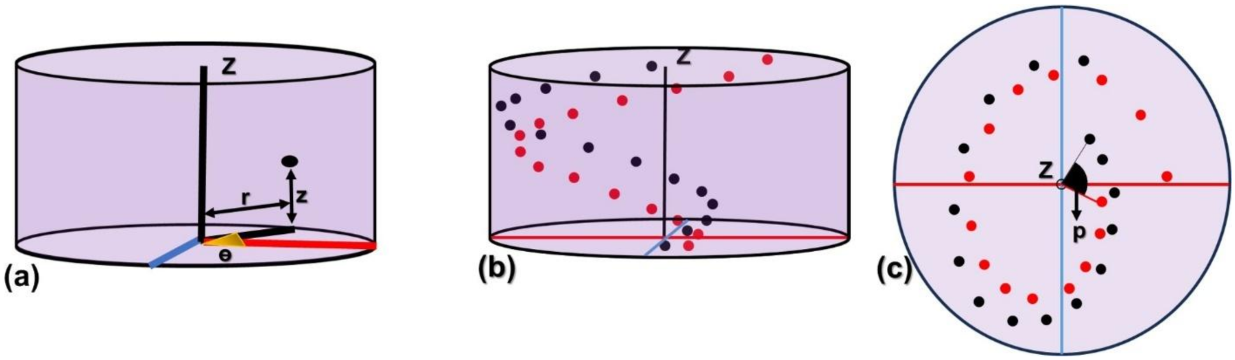

Figure 6: Presentation of the series of points in spiral axes to form the layers and segments in the cylindrical coordinate system. (a) The description of radius (r), azimuth (ɵ), and vertical distance (z) to define a point in the cylindrical coordinate system. The red line represents the radial axis, and the blue line represents a line perpendicular to it. (b, c) Longitudinal and transverse views, respectively, of the two series of points defined by the black and red dots in the cylindrical coordinate system, representing two consecutive layers/segments. Here, "p" marks the angular separation between the two series of points.

Values of r, ɵ, and z are generated so that the lamellae fully occupy the space, forming a compact skeleton in visual accordance with the 3D reconstruction obtained from the PXCT data. Here, the first and last points represent the base and the apex of the skeleton, respectively. Considering the total number of points in the series is n, an i^{th} point (r_i, $ɵ_i$, z_i), [where i<n] in the series is represented by the following equations:

$r_i = R + i*r_{step}$ [if i<=2*n/3] (radius increases for the first 2*n/3 points) and $r_i = r_{(i-1)} - i*r_{step}$ [if i>2*n/3] (radius decreases the last n/3 points) (1). This results in a globular shape for the skeleton.

R = Initial radius, r_{step} = (Maximum radius-Minimum radius)/n.

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$\Theta_i = (\Theta + i*\Theta_{step})$ (2)

Θ = Initial azimuth, Θ_{step} = angular separation between two consecutive lamellae = (Final azimuth-Initial azimuth)/n

$z_i = (Z + i*z_{step})$ (3)

Z = Initial vertical distance, z_{step} = (Final vertical distance-Initial vertical distance)/n

This results in a series of points with individual parameters, given by: [$(r_1, \theta_1, z_1), (r_2, \theta_2, z_2), \ldots.. (r_i, \theta_i, z_i), \ldots.. (r_n, \theta_n, z_n)$].

The difference between the final and the initial azimuths of the points of the series, is expressed as the angle of rotation (S). As a series of points is equivalent to a layer/segment of the skeleton, the angle of rotation of the series also represents the angle of rotation of a layer/segment. This angle of rotation (S) of a series/layer, is given by:

$S = \Theta_{step}*n$ (4)

A set of spiral axes (i.e., series of points), each with same numbers (n) of points is needed to create the other layers/segments for the entire skeleton reconstruction. Two such consecutive series of points (in red and black colors respectively) are presented in Figs. 6b and 6c. The points (r, θ, z) of all consecutive series is defined following the Eqs. (1), (2), and (3). The initial radius (R) and the vertical distance (V) is same for all the consecutive series of points. However, the initial azimuth (Θ) is different for each of the series and depends on the angular separation between two consecutive series of points. If the angular separation between two consecutive series of points is given by p. Then, the value of the p can be calculated as:

$p = 360°/N$ (5)

where N is the total number of series (i.e., the total number of layers/segments) for the full skeletal reconstruction. As the azimuth of the first point in the first series is θ_1, therefore the azimuth of the first point of the second series will be (θ_1+p). The I^{th} series of points in N numbers of series (I<N) can be represented as:

Series-I: [$(r_1, (\theta_1+(I-1)*p), z_1), (r_2, (\theta_2+(I-1)*p), z_2), \ldots.. (r_i, (\theta_i+(I-1)*p), z_i), \ldots.. (r_n, (\theta_n+(I-1)*p), z_n)$].

4.3.3 Creation of a layer and a segment

The segmented lamella is assigned to each point in the series, creating the layer and segment composed of lamellae. The inclination of the lamella is added by rotating it, as described in Figs. 5a and 5b. In the case of forming a layer (following model-1 of the microstructural arrangement), all the lamellae (n) are assigned a single value of angle of inclination (δ_A or δ_B). This suggests that rotating all the lamellae of the series with an inclination of δ_A or δ_B formed layer-A or layer-B, respectively. For generating a segment (following model-2 of the microstructural arrangement), the lamellae are alternatingly assigned two distinct angles (δ_A and δ_B) of inclination, indicating rotation of the alternative lamellae of the series with inclinations of δ_A and δ_B.

Considering that the first and the last points of the layer/segment represent the base and the apex of the skeleton, the tilt associated with the two points are τ_b and τ_a, respectively. The tilt values consistently change from τ_b at the base to τ_a at the apex. Thus, the value of tilt for ith point (i<n) of the series (i.e., layer/segment) is given by the following equation:

$\tau_i = \tau_{(i-1)}$ +i*tilt-increment (6), tilt-increment = $(\tau_b-\tau_a)/n$.

The value of the tilt is given to each lamella in the layer/segment by rotating it as described in Figs. 5a and 5c.

The descriptions of all parameters lead to the next step of reconstructing the full skeleton. The two skeletal reconstruction models of the microstructural arrangement have been termed as the layer model and segment model respectively. It is described earlier (in sections 2.2.1 and 2.2.2) that the total number (N) of layers/segments in the whole skeleton of *N. globulus* is calculated as 12. Thus, the angular separation between two consecutive layers/segments will, therefore, be p = 360°/12 = 30° (from Eq. (5)). However, the values of initial and final azimuths, initial radius and vertical distance, r_{step}, z_{step}, and the total number of lamellae (n) in a layer/segment are impossible to calculate without the segmentation of a layer/segment. As it is really difficult to separate one layer/segment of lamellae from the skeleton, we have applied several trials with different combinations of these values and have taken those values that give a space-filling layer/segment with the separated lamella. The results suggest that the total number of lamellae in one layer/segment (n) is 15. All the values of the parameters used to reconstruct the layers/segments, following different equations given in section 4.3.2 are given in Table 1.

Table 1: Values of the parameters required for reconstructing layers and segments of *N. globulus*.

Parameters	Values for layer model	Values for segment model
δ_A	25°	25°
δ_B	40°	40°
τ_a	-35°	-35°
Tilt-increment	4.37°	4.37°
τ_b	30.38°	30.38°
θ	0°	0°
θ_{step}	36°	6°
p	30°	30°
R	0.005 µm	0.005 µm
r_{step}	0.139 µm	0.139 µm

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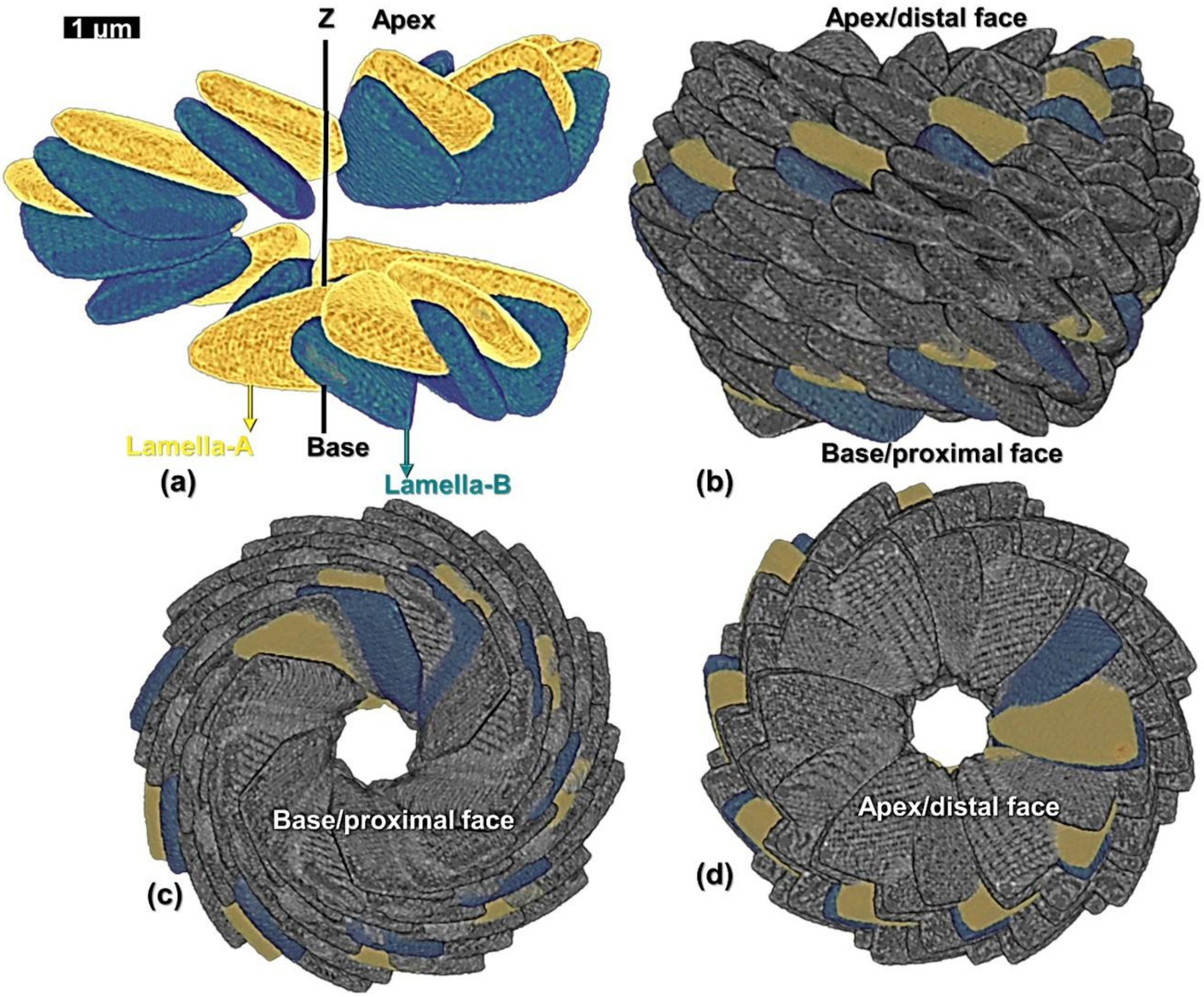
Z	0 µm	0 µm
z_{step}	0.251 µm	0.251 µm
n	15	15
N	12	12

4.3.4 Layer Model

Values of each of the cylindrical coordinates (r_i, θ_i, z_i), inclinations (δ_A and δ_B), and tilt (τ_i) for individual lamella in the first layer (layer-A) are provided in Table 2. The relative changes of these values for each of the lamellae across the length (L) are graphically represented in Fig. B4 in the Appendix. A total of 12 layers (six layer-A and six layer-B) are created, alternatively putting I = 0 to I = 11 in series I and using the values of parameters (layer model) of Table 1. Two successive layers of -A and -B and the full skeletal reconstruction are shown in Figs. 7a-7d.

Table 2: Values of the radius, azimuth, vertical distance, inclination, and tilt of all the 15 lamellae-A of the first reconstructed layer-A of *N. globulus*. The lamellae are labelled sequentially from A1 to A15.

Lamellae	Radius (r) (µm)	Azimuth (θ) (degree)	Vertical distance (z) (µm)	Inclination (δ) (degree)	Tilt (τ) (degree)
A1	2.09	0	0.47	20	-35
A2	2.23	36	0.72	20	-30.33
A3	2.37	72	0.98	20	-25.66
A4	2.51	108	1.23	20	-20.99
A5	2.65	144	1.48	20	-16.32
A6	2.79	180	1.73	20	-11.65
A7	2.93	216	1.98	20	-6.98
A8	3.07	252	2.23	20	-2.31
A9	3.21	288	2.48	20	2.36
A10	3.35	324	2.73	20	7.03
A11	3.49	360	2.98	20	11.7
A12	3.35	396	3.23	20	16.37
A13	3.21	432	3.49	20	21.04
A14	3.07	468	3.74	20	25.71
A15	2.93	504	3.99	20	30.38



305 **Figure 7: Reconstruction of the skeleton of *N. globulus* using the layer model. (a) Longitudinal view of the two layers formed by lamella-A (yellow) and -B (green). (b, c, and d) Longitudinal, basal and apical views, respectively, of the combination of 12 such reconstructed layers forming the entire skeleton. The first two layers of lamellae-A and -B are also shown. "Proximal" and "distal" (Aubry, 2013) terms refer to the orientation of the skeleton on the cell. Proximal face: face closest to the cell and distal face: face farthest from the cell.**

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4.3.5 Segment Model

Values of each of the cylindrical coordinates (r_i, $ɵ_i$, z_i), inclination (δ_A and δ_B), and tilt (τ_i) for individual lamella of the first reconstructed segment are given in Table 3. The graphical representation of their relative changes across the length (L) are presented in Fig. B5 in the Appendix. A total of 12 segments were generated using I = 0 to I = 11 in series I and the values of parameters (segment model) of Table 1. A segment with the lamella -A and -B and the full skeletal reconstruction are shown in Figs. 8a-8d.

Table 3: Values of the radius, azimuth, vertical distance, inclination, and tilt of all the 15 lamellae (eight lamellae-A and seven lamellae-B) of the first reconstructed segment of *N. globulus*. The lamellae-A and -B of the segments are sequentially marked from A1 to A8 and B1 to B7 respectively.

Lamellae	Radius (r) (µm)	Azimuth (ɵ) (degree)	Vertical distance (z) (µm)	Inclination (δ) (degree)	Tilt (τ) (degree)
A1	2.09	0	0.47	20	-35
B1	2.23	36	0.72	20	-30.33
A2	2.37	72	0.98	20	-25.66
B2	2.51	108	1.23	20	-20.99
A3	2.65	144	1.48	20	-16.32
B3	2.79	180	1.73	20	-11.65
A4	2.93	216	1.98	20	-6.98
B4	3.07	252	2.23	20	-2.31
A5	3.21	288	2.48	20	2.36
B5	3.35	324	2.73	20	7.03
A6	3.49	360	2.98	20	11.7
B6	3.35	396	3.23	20	16.37
A7	3.21	432	3.49	20	21.04
B7	3.07	468	3.74	20	25.71
A8	2.93	504	3.99	20	30.38

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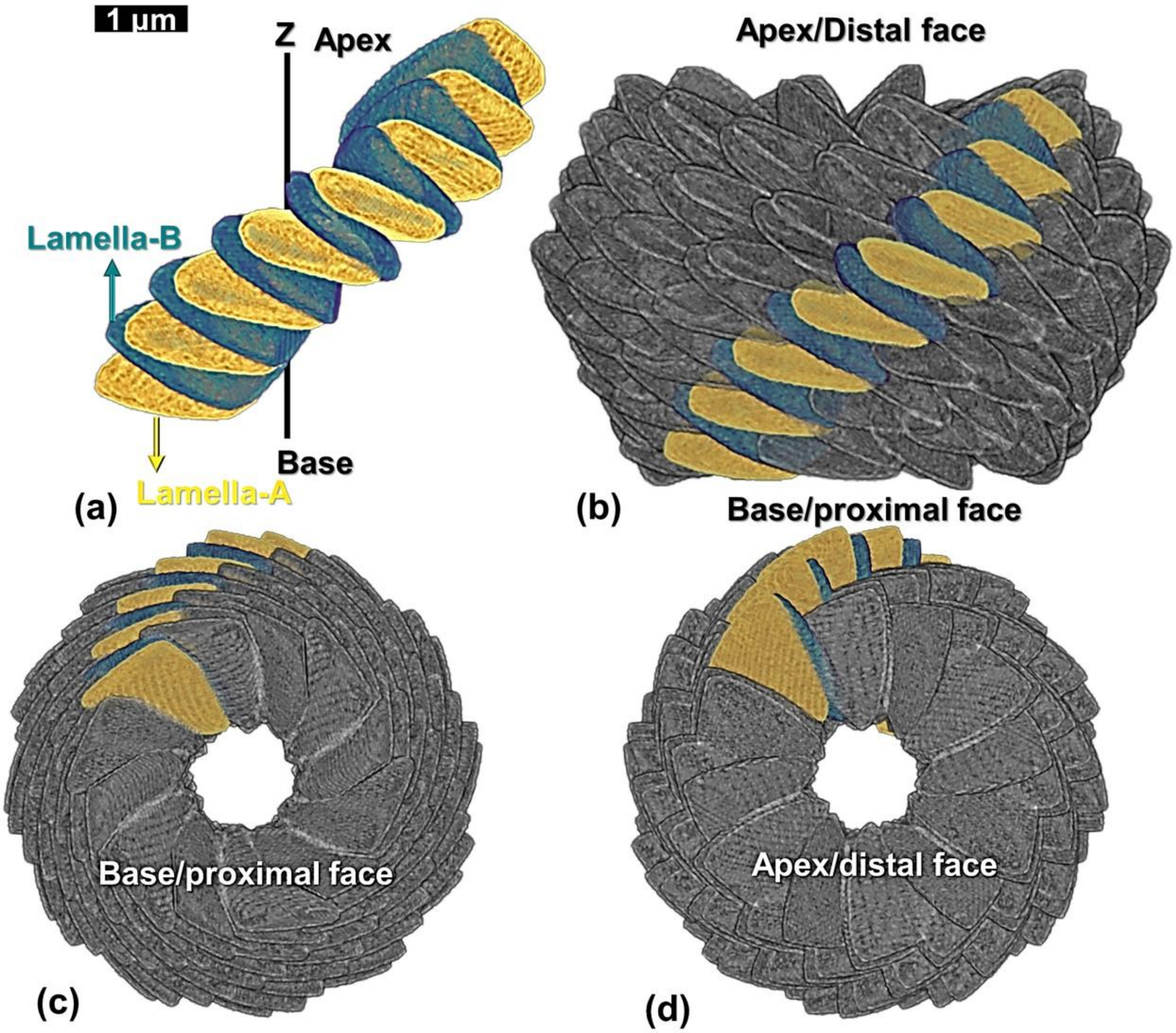

320 **Figure 8: Reconstruction of the skeleton of *N. globulus* using the segment model. (a) Longitudinal view with alternating, properly oriented, lamella-A (yellow) and -B (green). (b, c, and d) Longitudinal, basal and apical views, respectively, of the combination of 12 such reconstructed segments forming the entire skeleton. A segment containing alternatively placed lamella-A and -B is highlighted. "Proximal" and "distal" (Aubry, 2013) terms refer to the orientation of the skeleton on the cell. Proximal face: face closest to the cell and distal face: face farthest from the cell.**

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5 Discussion

The two models are based on two different concepts of skeletal microstructure of *N. globulus*. Both models result in similar reconstructions (Figs. 9a and 9b) of the skeleton, which structurally resembles the real specimen (Fig. 9c). However, each of these two models of the microstructural arrangement suggests a distinct biomineralization process to form the entire skeleton. The first model implies that the skeleton is formed by lamellae organized in successive layers, whereas the second one involves that the skeleton is formed primarily of segments, each formed by the stacking of lamellae. The structure "lamellae stacked in segments" (Aubry, 2013) as proposed in the second model is visible in *Braarudosphaera bigelowii*, an extant species with fossil representatives first occurring 100 Myr ago and belonging to the Order of *Braarudosphaerales* (Aubry, 2013) as *Nannoconus*. *B. bigelowii* calcifies a pentagonal skeleton with the combination of five trapezoidal segments (Fig. 4 in Hagino et al., 2016) composed of lamellae stacked in parallel to each other.

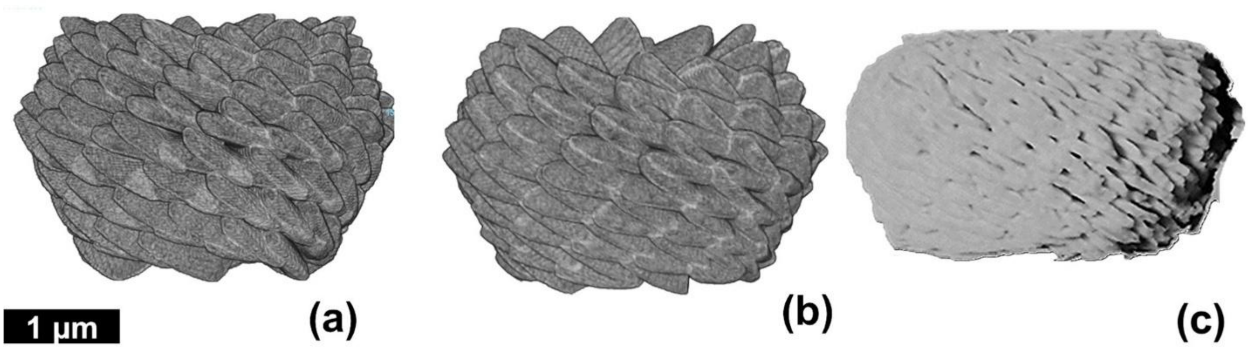

Figure 9: Two different reconstructions of the entire skeleton of *N. globulus*. (a) reconstruction based on the layer model and (b) reconstruction based on the segment model. Similar skeletal reconstruction is also proposed by Aubry (2025); Fig. 12d. (c) External view of the specimen of *N. globulus* used in the PXCT experiment (see also Fig. 3b).

Considering the segment model, we now try to show which parameters control the inter-specific morphological variability observed within the *Nannoconus* group during the ~30 Myr of the Early Cretaceous. The parameters that can potentially control the morphology of the skeleton are the angle of rotation of the segment and the radius. The angle of rotation of each segment (Eq. (4) in section 4.3.2) is considered as 90° in the discussed reconstruction by the segment model. With the same numbers of total segments ($N = 12$) and total lamellae in each segment ($n = 15$) used in the segment model, we have applied three different angles of rotations to individual segments: 30°, 60° and 180°. All of the angles of rotation used, resulted in reconstructed skeletons (Figs. 10a - 10d) of similar shape. So, it can be inferred that the angle of rotation of the segment does not affect the general shape of *Nannoconus*.

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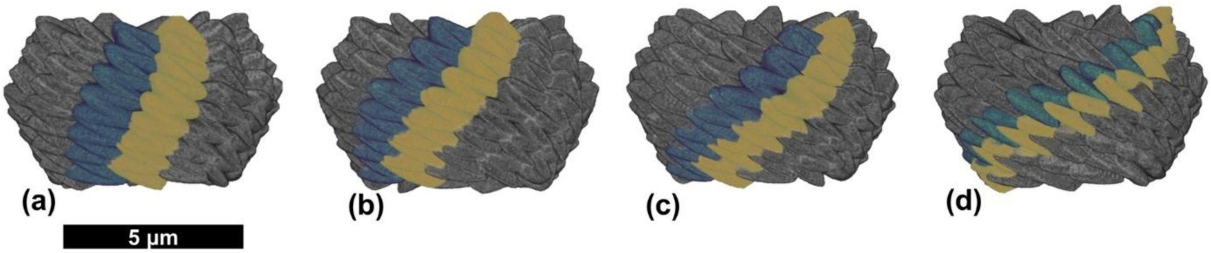

Figure 10: Four complete reconstructed *Nannoconus* skeletons based on the segment model with four different angles of rotation of each segment. (a-d) These angles of rotation are are 30°, 60°, 90°, and 180° respectively. Two consecutive segments of each reconstructed skeleton are marked in green and yellow.

The other parameter that can potentially affect the morphology of the skeleton is the radius. As we discussed earlier, the particular way of changing the radius with the length (see Eq. (1)) resulted in the globular shape of the skeleton. Thus, different ways of changing the radius would give rise to different skeleton's morphologies. Five different morphologies (globular, hour-glass, conical, barrel-shaped, cylindrical; Fig. 11) of the skeleton have been reconstructed by applying the same segment model using for each, the same values of the different parameters (Table 1) except for the radius (Table 4); they resemble the morphologies of *Nannoconus* real species. Thus, it is determined that the radius is the parameter that controls the inter-specific morphological variability of *Nannoconus*. However, other than radius, the total number of the segments forming the entire skeleton could also vary for different species of *Nannoconus*. It should be kept in mind that in this discussion the number of segments is considered to be 12 based on a SEM image of only one species, that is *N. globulus*. A comparative study of SEM images of different species of *Nannoconus* could be useful to investigate if the number of the segments in the skeleton is species dependent. More generally, with the model of reconstruction described in this study, we have succeeded in creating a 3D skeleton of *Nannoconus*; that means that we could constrain changing skeleton's morphology not only at the level of the Genus, but also at the level of the Order of *Braarudosphaerales*, allowing us to investigate whether a particular skeleton morphology had been favoured during the long evolutionary history (~150 Ma) of this order. Recent research (Aubry, 2025) suggests that the importance of this Order lies in its distinct skeletal structure, indicating both a taxonomic specificity and a unique biological position within the calcareous nannoplankton group.

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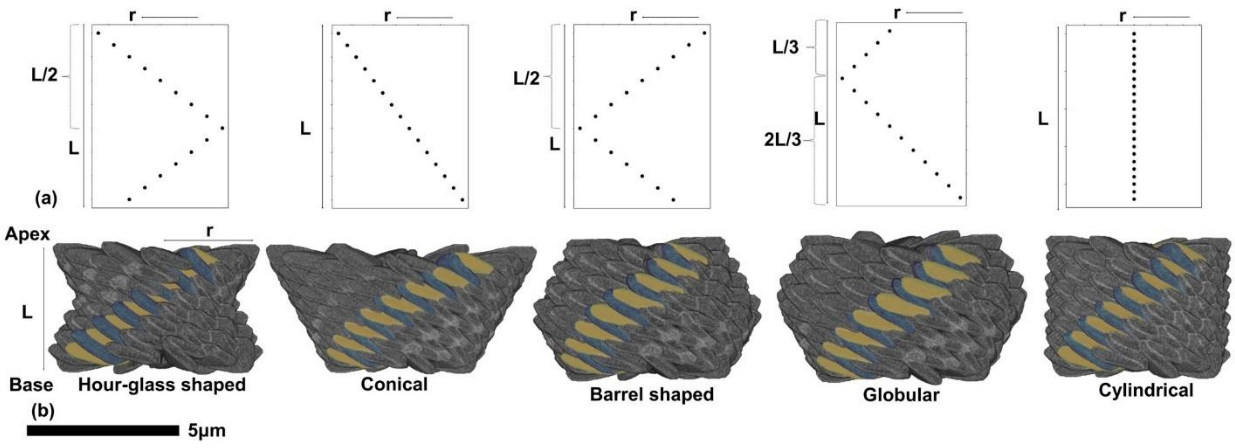

Figure 11: Five different skeletal morphologies of *Nannoconus*, each with distinct variation of radius (r) across the length (L). (a)
370 **Graphical expressions of five different variations of radius along the length of the reconstructed skeleton, as shown in Table 4. Each of the points on the graphs indicate individual lamella of a segment. (b) Skeletons reconstructed following each of the possibilities, using the segment model, with each skeleton highlighting a segment containing both types of lamellae-A (yellow) and -B (green).**

Table 4: Five possible variations in radius across the length of the reconstructed skeleton of *Nannoconus*. Each is associated with
375 **distinct skeletal morphologies (as shown in Fig. 11) and the corresponding species of *Nannoconus* exhibit these respective morphologies.**

Change of radius	Final morphology	Species
Radius decreases until ½ of the total length from the base and then increases to the apex	Hour-glass shaped	*N. abundans*
Radius constantly increases from the base to the apex	Conical	*N. steinmannii*
Radius increases until ½ of the total length from the base and then decreases to the apex	Barrel shaped	*N. circularis*
Radius increases until 2/3rd of the total length from the base and then decreases to the apex	Globular	*N. globulus*
Radius is constant from the base to the apex	Cylindrical	*N. truittii*

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It is now important to infer the process of biomineralization of the segments of the *Nannoconus*. Inferences can be provided based on the comparison of the biomineralization process of *B. bigelowii*, as documented in Hagino et al., (2016). In an un-calcified living cell of *B. bigelowii* an organic substrate is reported that resembles the shape of the pentagonal skeleton. The organic substrate is divided into five parts, with each of the parts mimicking the shape of a segment. In the vertical segments of the calcified skeleton produced by living *B. bigelowii*, thin organic layers are observed between two successive lamellae. Thus, the organic substrate and layers are seen to act as a "template" for the calcification of the segments and lamellae. This would indicate that the lamellae are stacked in each of the five parts of the organic substrate, templated by the organic layers, creating five segments. We suggest a similar process of calcification for *Nannoconus* with 12 segments; the calcification occurred on an organic "substrate" divided into 12 parts (since 12 segments are considered). Additionally, lamellae in the segments of *Nannoconus* are placed with inclination and tilt creating their interlocking arrangement whereas they are parallel in the segments of *B. bigelowii*. The inclination and tilt are explained by two mutually perpendicular rotations of the lamellae (see section 4.3.1) of *Nannoconus*. Such rotations with specific directions (i.e., clockwise and anti-clockwise) are commonly observed in biomineralized skeletons produced by marine organisms; for example, the coiling of gastropod's shell (Ueshima and Asami, 2003), the overlapping of the chambers generated by foraminifera (Schiebel and Hemleben, 2017), microstructural imbrications of the calcite units in the skeletons produced by calcareous nannoplankton (Young et al., 1999; Aubry, 2013, 2025). It is implied that these rotations are influenced by biomolecules such as the proteins, amino acids and polysaccharides (Young and Henriksen, 2003; Yu et al., 2005; Jiang et al., 2017, 2018, 2019). Was it the case for *Nannoconus*? To discuss this point, we presented here the process of mineralization of vaterite (a polymorph of calcium carbonate, $CaCO_3$) induced by amino acid (Jiang et al., 2018). The growth occurs layer-by-layer with amino acid intervening between two consecutive layers to form the microstructure of the layered vaterite. The two layers are termed "mother" and "daughter" layers. The amino acid (i.e., Aspartic acid) present between the mother and the daughter layers, rotates the daughter layer clockwise (Figs. 6A and 6B in Jiang et al., 2018) creating a difference in inclination between the two layers. Thus, because of the presence of amino acid between the two consecutive layers, the "inclination" of the new layers added during growth, increases continuously. The amino acid induced rotation of the layered vaterite structurally resembles the interlocking arrangement of the lamellae of *Nannoconus*. However, the reconstruction of the skeleton of *Nannoconus* showed that not only one but two rotations of the lamellae in two mutually perpendicular directions are necessary to develop the interlocking pattern (as discussed in section 4.3.1). Based on the comparison with the layer-by-layer grown vaterite, we hypothesize the probable presence of a layer of amino acid(s) or any biomolecule(s) to explain the two rotations between two successive lamellae of *Nannoconus*, creating the tilt and inclination. These successive lamellae could correspond respectively to the mother and daughter layers in the vaterite. For two such successive lamellae of a segment we call the preceding (similar to "mother layer" of Jiang et al., 2018) and succeeding (similar to "daughter layer" of Jiang et al., 2018) lamellae as the inferior and superior lamellae respectively (Fig. 12). The sense of the rotations to generate the inclination and tilt are marked in the inferior and superior lamellae in Fig. 12. The required arrangements for these two lamellae are: for inclination: opposite direction (both clock and anti-clockwise) of successive rotations (given by the black bold arrows in Fig.

12a), and, for tilt: the same direction (anti-clockwise) of successive rotations (given by the black dashed arrows in Fig. 12b). The opposite direction of rotations in the inferior and superior lamellae actually generate the two lamellar types i.e., lamella-A and -B. Therefore, both the clock and anti-clockwise rotations occur simultaneously during biomineralization of the superior lamella on the inferior lamella, generating the interlocking arrangement of the constituting lamellae. These possible explanations strengthen our initial hypothesis of the amino acid(s)/biomolecule(s) containing layer, between two successive lamellae. This layer, containing the amino acid(s)/biomolecule(s), could correspond to the organic substrate occurring as the "template" for the calcification of the segments of *Nannoconus*.

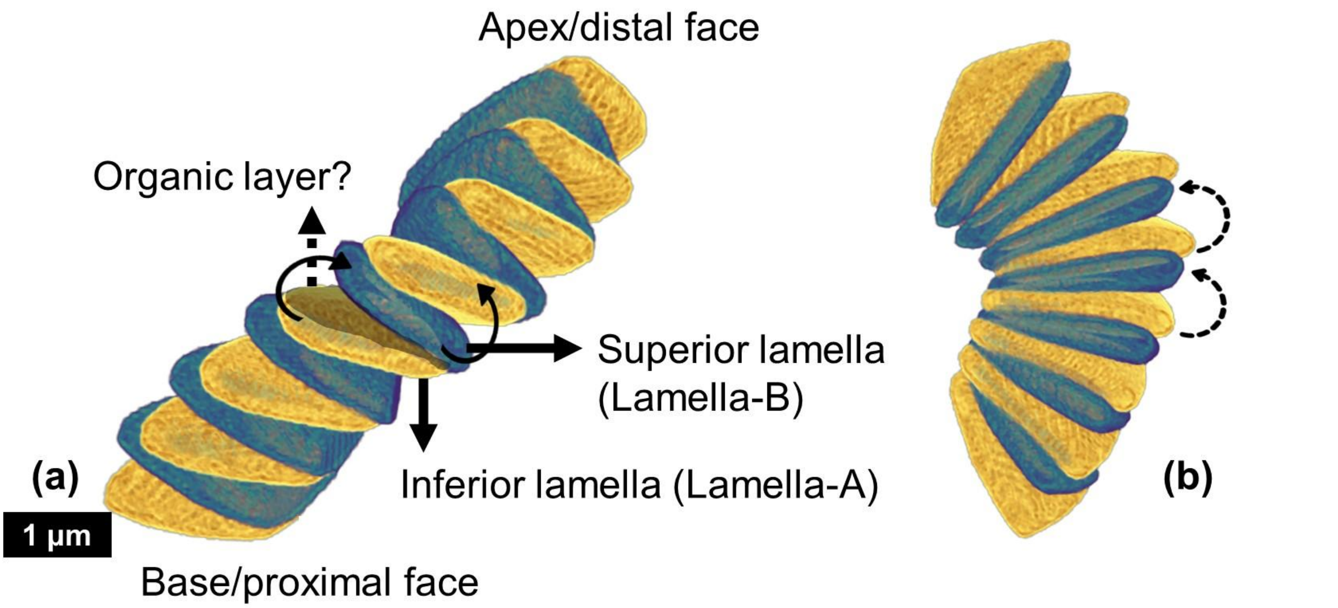

Figure 12: A reconstructed segment of *N. globulus* with the proposed inferior and superior lamellae along with the hypothesized intervening organic layer between them. (a) The two black bold arrows show the two opposite directions of rotation (clock and anti-clockwise), which create distinct inclinations in the two successive lamellae, generating the lamella-A (yellow) and -B (green). (b) Same segment, with two dashed black arrows presenting anti-clockwise rotations in two successive lamellae, generating the tilt in them.

These organic layers, if they were present in the *Nannoconus* skeleton, are unlikely to be preserved in the fossil record. However, specimens of *Nannoconus* have been studied with confocal Raman micro-spectroscopy, and the obtained spectra have shown the presence of preserved "organic matter" within the skeleton (Fig. B6 in the Appendix). The confocal Raman spectra of *Nannoconus* are characterized by a number of peaks, with distinct intensities at specific wavenumbers. The spectra present two distinct parts: one from 150 to 1200 cm^{-1} with characteristic peaks of the skeletal $CaCO_3$ and the second one from 1200 to 2000 cm^{-1} which could correspond to the "organic matter" signature preserved within the skeleton. Indeed, in a study focused on the identification of organic matter in bioclastic grains encountered in marine carbonate rocks, Moya et al., (2023) performed Raman micro-spectroscopy and detected on the spectra the fingerprints of organic matter in the higher wavenumber region (i.e., wavenumber>1200 cm^{-1}). Additionally, Raman signals related to protein and polysaccharide are also reported in the same spectral region from biocalcite produced by extant calcareous nannoplankton (Silvestri et al., 2020). Therefore, a future investigation using tomographic technique with resolution finer than the PXCT, such as atom probe tomography (resolution ~0.5 nm) could be useful to confirm the presence of organic matter preserved within the skeleton of *Nannoconus*.

We have hypothesized above that the biomineralization of the lamellae of the *Nannoconus* segments is "templated" by an organic layer containing amino acid(s)/biomolecule(s). The rotations induced by the amino acid(s)/biomolecule(s) created the inclination and tilt associated with the lamellae hence generating the interlocking arrangement of the lamellae. Both the clock and anti-clockwise rotations are inferred in the same skeleton of *Nannoconus*. Such "biomolecule(s)" influenced natural biomineralized 3D microstructure with repetitive arrangement of units like the calcite lamellae as observed in *Nannoconus*, are often studied to synthesize biomimetic materials (Jiang et al., 2019). These materials engineered for enhanced physical (e.g., hardness) and chemical (e.g., solubility) properties (Cho et al., 2023) have applications in research fields like catalysis and biomedicine. Therefore, the 3D skeleton reconstruction of *Nannoconus* can definitely provide significant insights to biomimetic material design, expanding its relevance beyond Paleontology.

6 Conclusion

Among calcareous nannofossils, the *Nannoconus* group of the Order *Braarudosphaerales* (fossil and extant representatives) presents sophisticated massive skeletons, organized in an interlocking arrangement of calcitic lamellae stacked in segments, forming a wall around a central canal. They contributed to huge carbonate accumulations over ~30 million years in the Early Cretaceous seas. The biomineralization process that led to the production of these massive skeletons was non-existent. The aim of the present study was a proper understanding of this process from a 3D reconstruction of the skeleton. A set of Ptychography X-ray computed tomography (PXCT) with synchrotron radiation at SWING Beamline of SOLEIL (French synchrotron) was applied on several well-preserved *Nannoconus* skeletons to understand the microstructural arrangement at the nanometer level (finer than the thickness of a lamella). The result of the experiment was a series of tomographic image slices (3D resolution ~40 nm) for the skeleton. Then, one lamella, virtually separated from the image slices, is used to reconstruct the entire skeleton of *Nannoconus* in the ORS-Dragonfly software. For the first time, a 3D skeletal reconstruction has been achieved for *Nannoconus globulus*. The interlocking arrangement of the lamellae results from two different angles termed inclination and tilt, while changing radius (minimum ~2.1 µm to maximum ~3.5 µm) controls the skeletal morphology. The successful 3D reconstruction of a skeleton using segmented lamellae of *Nannoconus*, validates the segment model, that is the skeleton is a spiral combination of identical, imbricated segments of stacked lamellae, as previously

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suggested by Aubry (2013). The 3D model of skeleton reconstruction developed in the present study could be further applied to the reconstruction of the skeleton of other Genera within the Order of *Braarudosphaerales* to investigate if a particular skeleton morphology has been favoured during the long evolutionary history of this order. The biomineralization of the segments is hypothesized to be templated by organic layers containing amino acid(s)/biomolecule(s). During the biomineralization of the lamellae, the amino acid(s)/biomolecule(s) is proposed to have induced clockwise and anti-clockwise rotations, creating the inclination and tilt and therefore generating the interlocking arrangement of the lamellae. Biomolecule(s)-driven naturally produced microstructures characterized by the repetition of units such as the 3D arrangement of the lamellae of *Nannoconus's* skeleton, are often extensively investigated for designing biomimetic materials. Such materials with artificially enhanced physico-chemical properties (e.g., hardness, solubility) are implemented in research with specialisation to catalysis and biomedicine, therefore making this publication relevant and important, outside the study of calcareous nannofossils.

Appendix

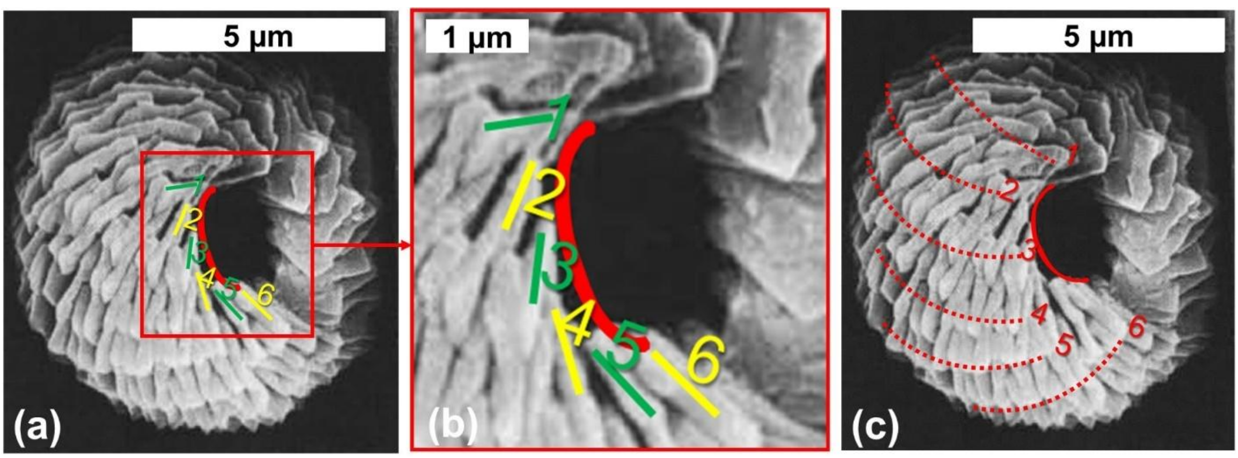

Figure B1: Three quarter view of the distal face of *N. globulus* in scanning electron microscopy (SEM). (a) Six lamellae in the 180° transect of the apex (given by the red curve) are highlighted. Three of lamellae-A and three of lamellae-B are numbered in green and yellow colours respectively. (b) Magnified view of the part delineated by red box in (a) with all the six lamellae: three lamellae-A (yellow) and three lamella-B (green). (c) Boundaries between six segments highlighted by red-dotted lines in the same 180° transect of the apex of the *N. globulus*. SEM image adapted from Covington and Wise (1987).

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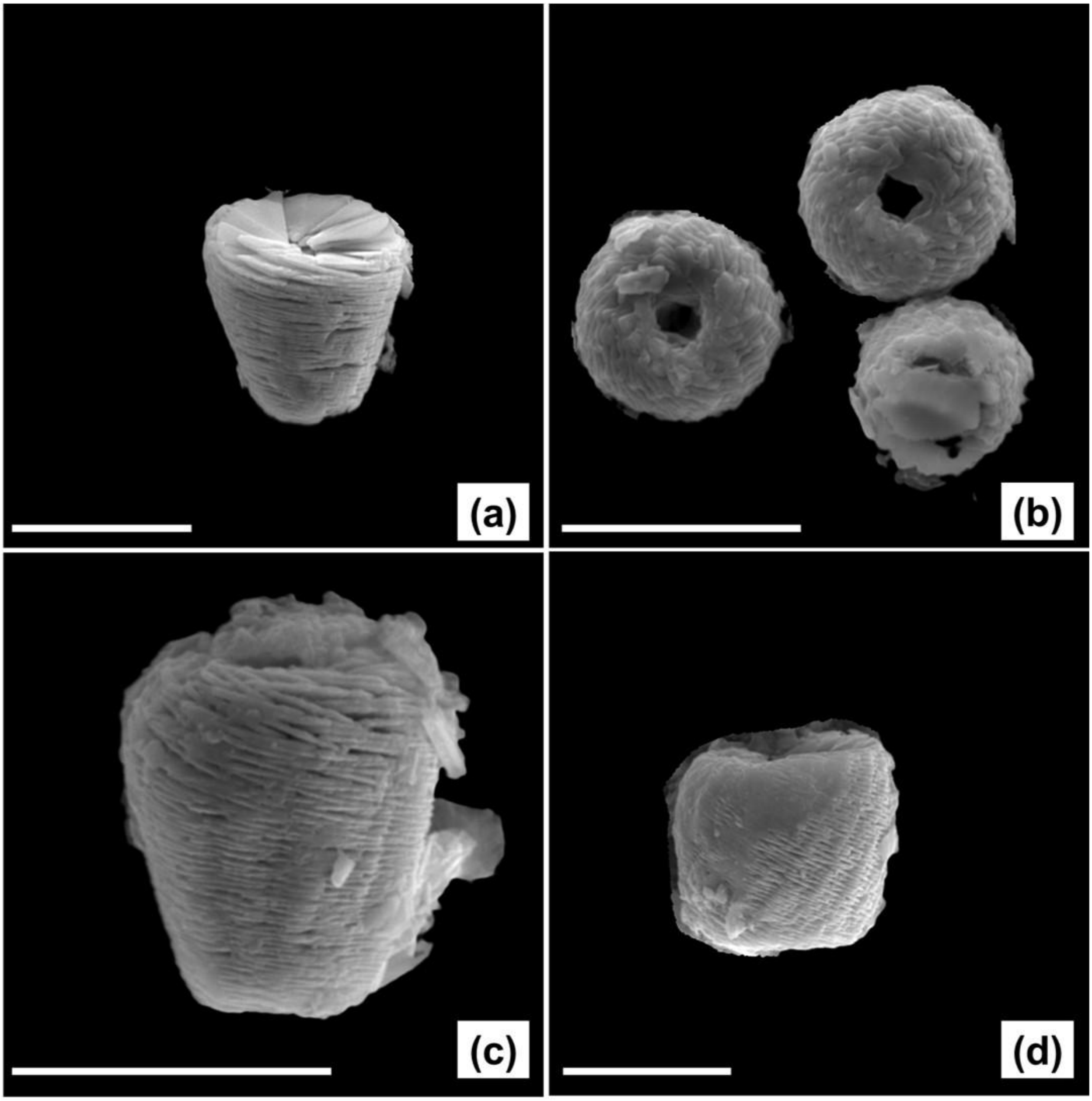

Figure B2: SEM images of well-preserved specimens of *Nannoconus*, from DSDP Leg-93-Site 603 (Atlantic Ocean, lower continental rise of Cape Hatteras) core 44, interval 115-116 cm. Individual lamella can be clearly distinguished in each of the specimens because of their good preservation. Identification: (a) *N. steinmannii* (b) *N. globulus* (c) *N. kamptneri* (d) *N. circularis*. The white line represents 10 µm.

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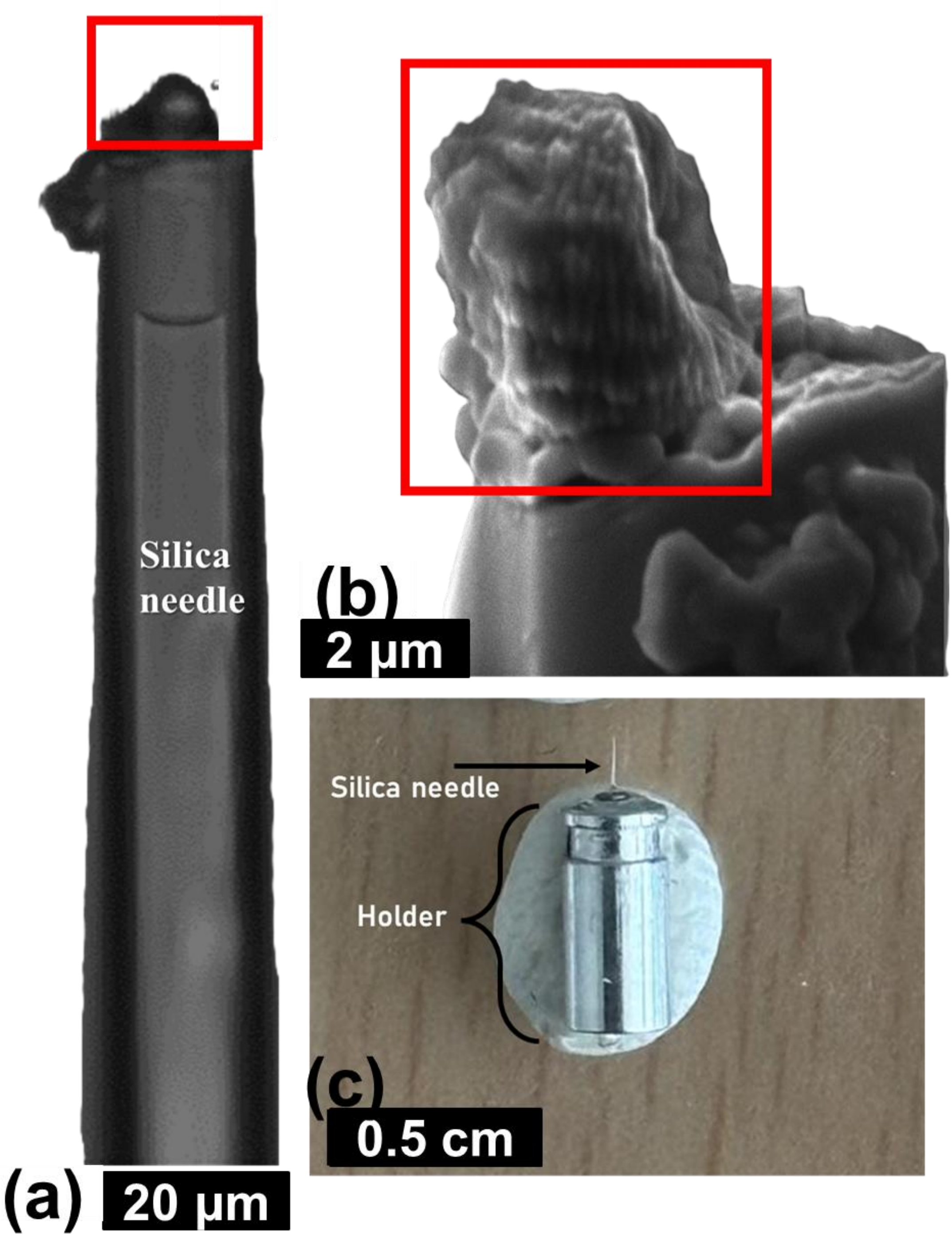

490 **Figure B3: Procedures used for the sample preparation of synchrotron-based ptychography X-ray computed tomography experiment, applied on *Nannoconus globulus*. (a) A specimen of *N. globulus* (inside the red box) is manually picked with a silica needle. (b) SEM image of the specimen. (c) The needle with a *Nannoconus* resting at its tip is placed in a metal holder that fits to the experimental station of the beamline of the synchrotron radiation centre.**

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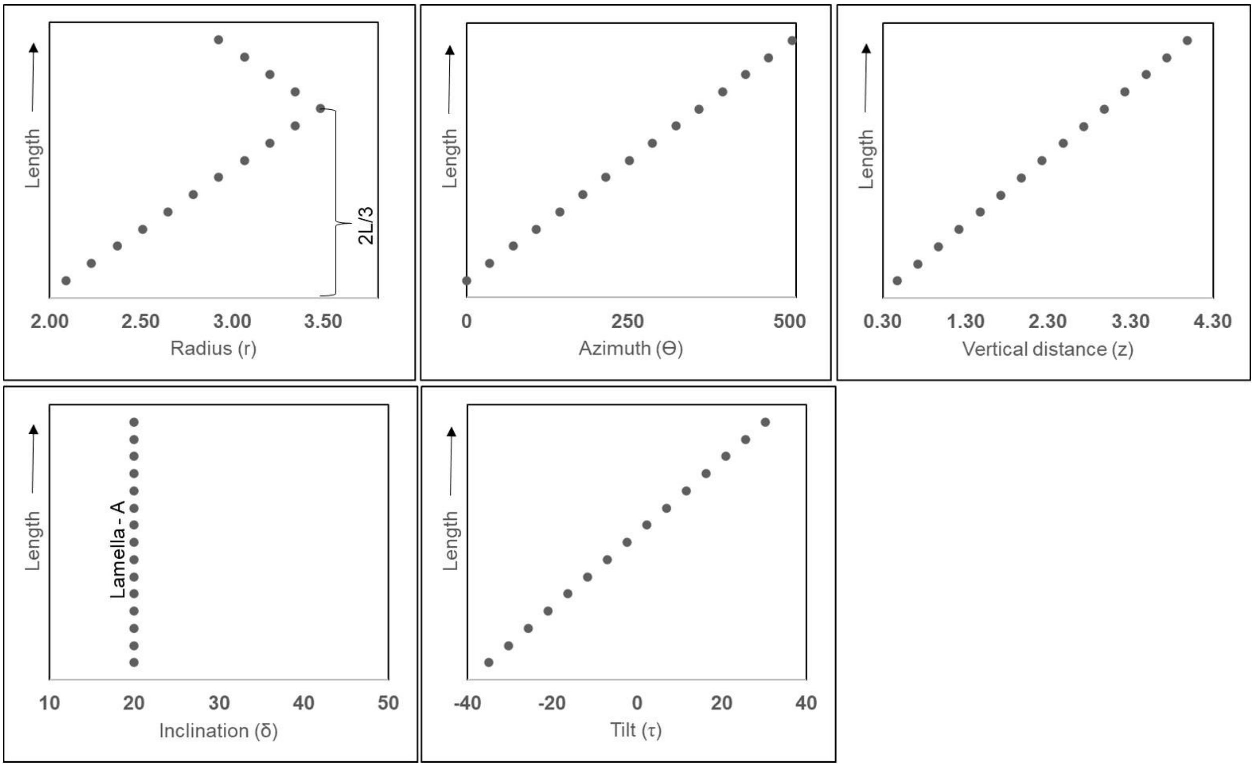

Figure B4: Graphical representation showing the variation of parameters along the length (L) of the first reconstructed layer (layer-A) composed of lamella-A, of *N. globulus*. The black points in the graphs, mark individual lamellae.

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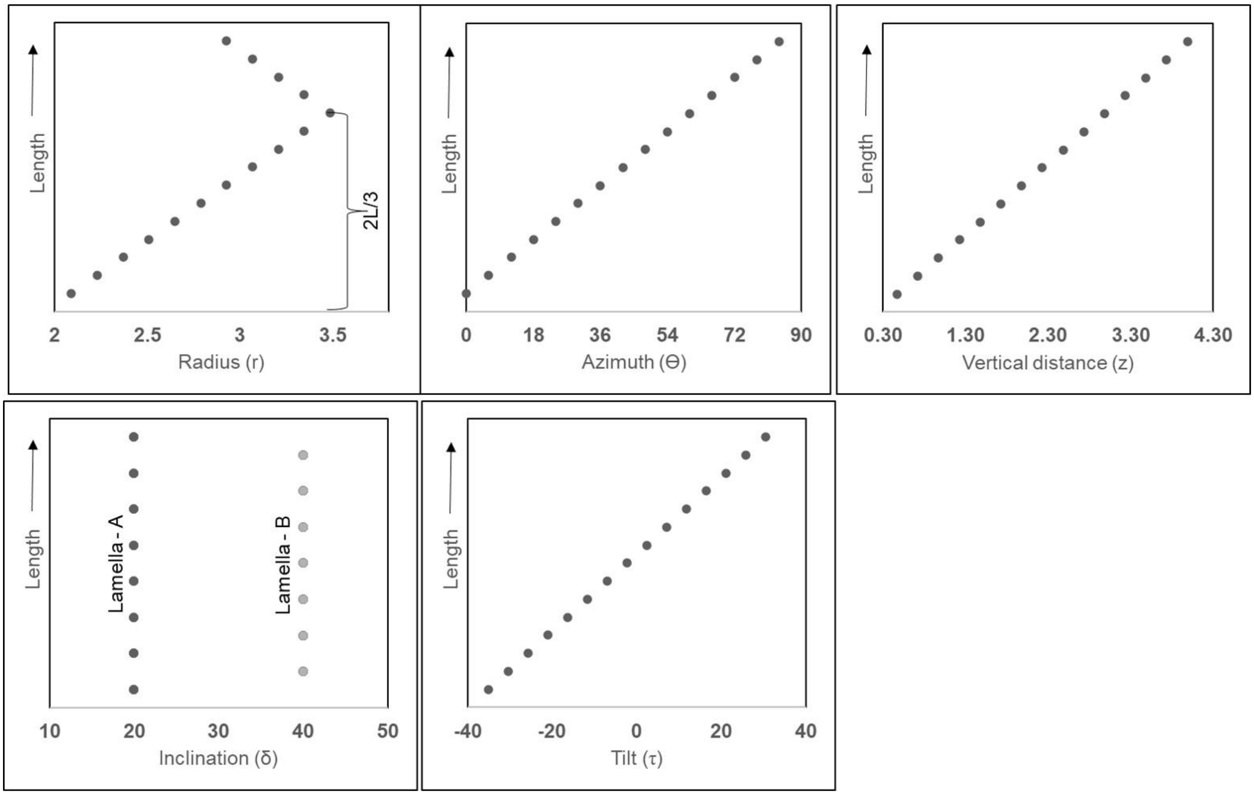

Figure B5: Graphical representation showing the variation of parameters along the length (L) of the first reconstructed segments composed of lamella-A and -B of *N. globulus*. The black points in the graphs, mark individual lamellae.

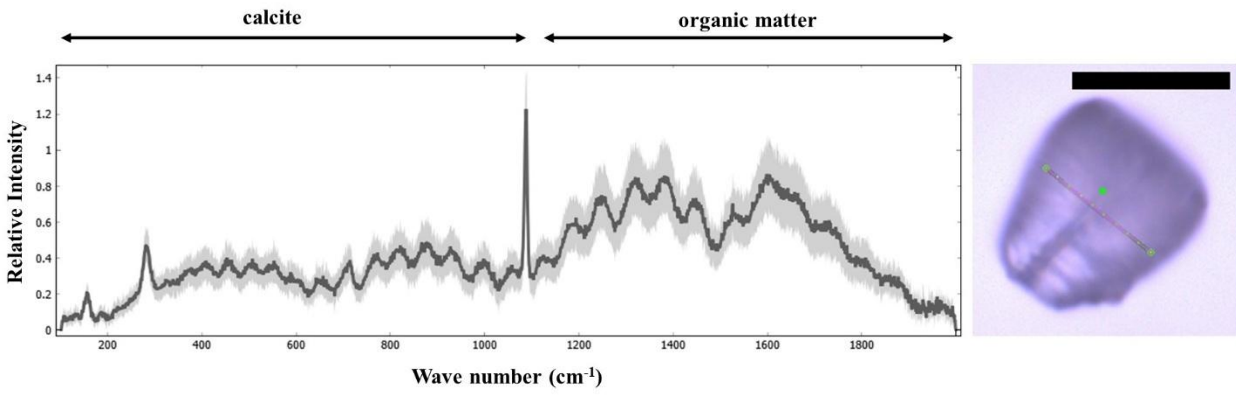

Figure B6: Confocal Raman spectra of *Nannoconus* displaying two regions corresponding to the compositional $CaCO_3$ and the "organic matter" preserved within the skeleton. The photomicrograph in optical microscopy (PPL) of the *Nannoconus* is given on the right, with a red line showing the scanning path for the spectra. The black scale bar represents 10 µm. PPL: plane-polarized light. The Raman spectra were obtained using a LabRAM Soleil Horiba Raman micro spectrometer with 532 nm wavelength laser-excitation and 1 µm spot size. A laser power of 1.1 mW was used to confirm that there is no damage of the specimens induced by the laser. The laser beam was focused on the sample through a hole of 50 µm using an objective at 100X magnification. The specimen was subjected to linear scans along the surface (5-10 µm) and also inside the specimen (4-5 µm). Total 25 scans were recorded, each with an acquisition time of 2 seconds. Peaks were fitted in the spectrum using Gaussian functions in Quasar

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software (Toplak et al., 2017) as a part of the analyses. The spectra have been normalized to the maximum intensity. The black spectrum represents the average of all 25 recorded spectra of the *Nannoconus*.

Table C1: Glossary of terms used in this work.

Term	Description
Taxonomy	A formal hierarchical biological classification scheme of any organism, also known as Linnaean Taxonomy. This is given by Life<Domain<Kingdom<Phylum<Class<Order<Family<Genus<Species
Order	The 6^{th} rank of the taxonomic classification.
Genus	The 8^{th} rank in the taxonomic classification, the plural of genus is genera.
Species	The 9^{th} and the final rank of the taxonomic classification.
Morphogroups	An informal term to classify organisms based on similar morphological criteria.
Convex Hull	A mathematical representation of an envelope/elastic cover covering a large set of data, defining a shape in space.

Code availability

The code for the skeletal reconstruction of *Nannoconus globulus* for both the layer and segment models is available in the repository: https://doi.org/10.5281/zenodo.14925063 (Chowdhury et al., 2025)

The segmentation, reconstruction, and visualization of the results have been done in Dragonfly software version 2024.2 for [windows]. Comet Technologies Canada Inc., Montreal, Canada; software available at https://www.theobjects.com/dragonfly.

Sample availability

Rock samples and associated nannofossil samples used for the experiment are curated at the Collections de Géologie de l'Observatoire des Sciences de l'Univers de Grenoble (OSUG), with an appropriate UJF-ID number. OSUG-COLLECTIONS is a database of rocks, minerals, and fossils, https://web.collections.osug.fr, OSUG, UGA. doi:10.17178/OSUG-COLLECTIONS.all.

Author contributions

RC prepared the sample with inputs from **BSM**, **FG**, **AK,** and **AFM**, applied the image segmentation, co-developed the reconstruction code with **AFM,** and wrote the original draft. **JCS**, **JLH,** and **AK** helped in the initial conceptualization of the experiment. **MPA** intensely contributed to the conceptualization of the segment model of the reconstruction. **RC**, **RB**, **BSM**, **MD**, **AK**, **JP**, **FG,** and **AFM** performed the experiments in the synchrotron radiation centre. **FG** and **AFM** jointly supervised **RC** in this work and acquired all the relevant funding. All the co-authors contributed significant inputs for the review and editing.

Competing interests

The authors declare that they have no conflict of interest.

Acknowledgments

We acknowledge SOLEIL for providing access to synchrotron radiation on the SWING beamline through proposals 20211643 and 20221681.

Financial support

This work was supported by the Tellus Program of CNRS-INSU, OSUG@2020, and IODP-France.

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