The argonaut constructs its shell via physical self-organization and coordinated cell sensorial activity.
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SUMMARY
The shell of the cephalopod Argonauta consists of two layers of fibers that elongate perpendicular to the shell surfaces. Fibers have a high-Mg calcitic core sheathed by thin organic membranes (>100 nm) and configure a polygonal network in cross section. Their evolution has been studied by serial sectioning with electron microscopy-associated techniques. During growth, fibers with small cross-sectional areas shrink, whereas those with large sections widen. It is proposed that fibers evolve as an emulsion between the fluid precursors of both the mineral and organic phases. When polygons reach big cross-sectional areas, they become subdivided by new membranes. To explain both the continuation of the pattern and the subdivision process, the living cells from the mineralizing tissue must perform contact recognition of the previously formed pattern and subsequent secretion at sub-micron scale. Accordingly, the fabrication of the argonaut shell proceeds by physical self-organization together with direct cellular activity.

INTRODUCTION
The shells of proto- and metazoans are organomineral biocomposites consisting of recurrent arrangements of crystals and organic matrices, called microstructures (sometimes referred to as “structures” or “textures”) (e.g., Carter et al., 2012). Microstructures are adapted for their mainly protective function, although they also provide structural support for the soft parts, prevent dissolution, help in locomotion, etc. Their impressive biomechanical properties make them highly valued scientific objects for study. Accordingly, there is strong interest in knowing the strategies used by organisms for the production of microstructures to get inspiration for the fabrication of similar highly functional synthetic products.

Except for siliceous sponges, linguliformean phosphatic brachiopods, and chordates, metazoan skeletons are based on calcium carbonate (Lowenstam and Weiner, 1989). The different taxa also have different abilities for the production of microstructures (Checa, 2018). Mollusks are, by far, the leaders in both the variety and sophistication of their microstructures. They are able to produce some twelve basic microstructures with either aragonite or calcite. Both polymorphs can be present in a single shell, where the calcite layers, if any, are always external (Bagnold, 1930). Taxa such as brachiopods (Simonet Roda et al., 2021), bryozoans (Taylor et al., 2015), and polychaetes (Vinn, 2020) are also able to produce each a moderate suite of microstructures, sometimes with only a single mineral (e.g., calcite in articulate brachiopods). Corals (Stolarski, 2003) and cirripeds (Bourget, 1987) can only secrete one or two monomineralic microstructural types.

From a strict proteomic approach, proteins are responsible for every aspect of microstructures (Heywood, 1994). Studies show that soluble proteins control the mineral phase (aragonite, calcite) (Balcher et al., 1996; Falini et al., 1996), as well as nucleation and crystal shape (Marin, 2020). But some aspects of microstructures cannot be produced by the biomolecular toolkit directly. For example, some prismatic layers of mollusks consist of calcitic or aragonitic prisms surrounded by relatively thick organic membranes (Griesshaber et al., 2018). Nacre is made of an alternation of lamellae of tablets and thin organic membranes. These and other examples have been alternatively and successfully explained by self-organization: emulsification in the prismatic layers (Checa et al., 2016a) and layer-by-layer liquid crystallization in nacre (e.g., Cartwright and Checa, 2007), sometimes coupled with a particularly sophisticated activity of the mantle cells...
Within mollusks, cephalopods are a rather restricted group in terms of diversity (830 accepted extant species), fourth after gastropods (70,744), bivalves (9,731), and Polyplacophora (1,046) (WoRMS Editorial Board, 2021a, 2021b). Within mollusks, cephalopods are a rather restricted group in terms of diversity (830 accepted extant species), fourth after gastropods (70,744), bivalves (9,731), and Polyplacophora (1,046) (WoRMS Editorial Board, 2021a, 2021b). The female Argonauta (also called paper nautilus) secretes a thin spirally coiled shell, made entirely of calcite. The minute male is devoid of any shell. The first detailed observations on the secretion of the shell by the animal were made by Jeanne Villepreux-Power in the 19th century. In her (sometimes overlooked) pioneering study on Argonauta argo (Power, 1856), she described how the female argonaut shell is secreted by very wide membranaceous distal extensions (webs) of the two dorsal arms, and not by the mantle, as in all mollusks. This observation was confirmed by later authors (Finn, 2013; Milner, 1935; Mitchell et al., 1994; Woodward, 1880). Details about how shell growth by the webs proceeds are still lacking (Oudot et al., 2020). Accordingly, this shell is not homologous to the mollusk shell but an apomorphy of Argonauta. As might be anticipated, the shell of Argonauta is also unique in its general structure, and microstructure. Unlike the Nautilus, the female argonaut shell is devoid of septa, siphuncle, or any other internal structure, including attachment sites of the soft parts (Adams and Adams, 1858; Power, 1856; Woodward, 1880). It is used by the female to both brood the eggs (egg-case) and trap air inside for buoyancy purposes.

The first observations of the microstructure of the shell of Argonauta were done with optical microscopy by Bøggild (1930), who observed that the shell consists of two finely prismatic layers, which meet at a central dark layer. Later, with the employment of electron microscopy, Kobayashi (1971) confirmed Bøggild’s observations, revealed the organic nature of the central layer, and provided accurate details regarding the fibers making up the prismatic layers. Their diameters are less than one micron and consist of a calcitic core and an envelope of “reticulated” organic membranes. Later authors (Bandel and Dullo, 1984; Wolfe et al., 2012, 2013) overlooked Kobayashi’s model and concentrated on the calcitic phase of the fibrous bidirectional microstructure. Only Mitchell et al. (1994) showed interesting details regarding the organic network surrounding the calcitic fibers, which they qualified as polygonal, typical of minimal surface energy configuration.

We have carried out an in-depth study of the microstructure of the shell of Argonauta using high-resolution microscopy techniques. We focused particularly on the evolution of the organomineral complex during growth and considered if it can be explained by means of self-organization, cell activity, or any other phenomenon. The fact that the Argonauta shell is secreted by an organ different from the mantle might lead one to think that it must be a rudimentary construct. We found quite the contrary. Its constructional route is extremely sophisticated, similar, or even more sophisticated than those followed by other mollusks to produce similarly complex mantle-made materials.

**RESULTS**

**Mineralogy and composition**

X-ray diffraction (XRD) analysis indicates that calcite is the only mineral phase present. The results of the Rietveld analysis (Table S1) show well-defined changes in calcite unit cell parameters with respect to pure inorganic calcite, which can be explained by both the substitution of Ca\(^{2+}\) by Mg\(^{2+}\) ions and the incorporation of organic molecules into the calcite crystal structure (Pokroy et al., 2006). In particular, the reduction in calcite unit cell parameters observed in the sample treated at 400°C (Δa = −0.0432 ± 3 × 10\(^{-4}\) Å = −0.865%, Δc = −0.19678 ± 9 × 10\(^{-3}\) Å = −1.153%), when the influence of organic matter is negligible, can be explained by a 10.4% (molar) substitution of Ca\(^{2+}\) by smaller Mg\(^{2+}\) ions (Dos Santos et al., 2017). The expansion of calcite unit cell parameters (Δa = 0.107%, and Δc = 0.082%) observed in the untreated shell mineral compared with the sample treated at 400°C (no organic matter) may result from the incorporation of organic shell components into the calcite crystal structure.

The amount of amorphous material estimated by XRD using silicon as an internal standard was 6.88% by weight. On the other hand, thermogravimetry analysis coupled with differential scanning calorimetry
(TGA-DSC) revealed ~1.5% of structural water (weight loss up to 200°C) and 8.68% of organic matter (weight loss between 200% and 600%), that is, a total of 11.91% of amorphous material, notably higher than the amount calculated by XRD. This indicates that XRD may not be sensitive enough to quantify all amorphous components.

Fourier transform infrared (FTIR) spectra of the organic fraction (Figure S1) show strong protein peaks (Amide I and Amide II) and low-intensity absorption peaks in the polysaccharide region.

**Shell structure and microstructure by electron and atomic force microscopy**

Scanning electron microscopy (SEM) observations reveal that the shell consists of two finely prismatic or fibrous layers separated by a central organic layer (Figure 1A), as previously observed (Bøggild, 1930). At the boundaries of the central organic layer with both fibrous layers, there is a series of centers from which...
radiating arrangements of fibers (i.e., spherulites) initiate (Figures 1B–1D). Upon growth, they compete with each other, such that fibers become increasingly parallel (Figures 1D and 1E). Growth lines, marking the positions of the mineralizing tissue, extend uninterruptedly across the contiguous spherulites (Figures 1A, 1B, 1D, and 1E) and fibers (Figure 1F). Upon decalcification, the central layer reveals a high amount of organic substance and the material retains its whole spherulitic fibrous aspect (Figure 1G). In close view, the decalcified fibers consist of an organic network of membranes forming very elongated prismatic units with polygonal outlines (Figures 1H and 1I), as described (Kobayashi, 1971; Mitchell et al., 1994). Growth lines are imprinted on the surfaces of the membranes (Figure 1H, inset).

Transmission electron microscopy (TEM) examination provides additional details of the organic network. Sections perpendicular to the growth surfaces reveal that every spherulite has an initial dense organic-rich core, from which both the calcite fibers and the intervening membranes initiate (Figures 2A–2C). More to the interior of the central layer, the organic material looks looser. The boundaries between spherulites are marked by membranes (Figure 2D). Sections parallel to such surfaces reveal the reticulated pattern formed by the organic membranes (Figures 2E and 2F). It is formed by polygons with diameters between less than 100 nm to close to
2 μm. The number of sides ranges from three to eight, also depending on size. As previously indicated (Mitchell et al., 1994), the polygons resemble minimal surface structures whose sides meet at triple points, although quadruple points are not infrequent (Figure 2F). The organic pattern can be qualified as cellular, but we will avoid the term “cell” to refer to individual polygons in order to avoid confusion with the cells of the living tissue, referred to below. Frequently, membranes extend from one side toward the interior of the polygon and do not reach the opposite side (Figures 2E and 2F), something also observed under the SEM (Figure 1I, inset), and previously noticed (Mitchell et al., 1994). More rarely some short membranes appear isolated within single polygons (Figures 2E and 2F, inset). Membranes show consistent thicknesses between 70 and 90 nm, and are constituted by a double layer, separated by a distinct dark interphase a few nanometers thick (Figure 2G).

In transverse section, each layer consists of aligned rounded elements, which, in longitudinal section, appear noticeably elongated (Figures 2B, 2C, 2H, and 2I). All the above implies that the organic walls are made by two adjoining palisades of fibers elongated in parallel to the growth direction and separated by a dark interphase a few nanometers thick.

TEM and atomic force microscopy (AFM) observations reveal that the calcite constituting the shell of Argo-nauta has an ultrastructure similar to that of most other calcium carbonate biominerals. TEM reveals the presence of wide amorphous rims (sometimes >10 nm) around the crystalline phase (Figures 3A and 3B), which can be recrystallized by beam radiation (Figure 3B). This argues for a minority ACC phase. AFM observations (tapping mode) (Figures 3C and 3D) reveal a distinct surface roughness, typical of biocrystals (Dauphin, 2008), with two phases with different contrast (phase mode): a lighter contrast phase forming the bulk of the structure and a dark minority phase in the form of threads (arrows in the amplitude and phase images of Figure 3D). Previous studies on other biocrystals interpreted the lighter phase as crystalline and the darker phase as ACC enriched in biomolecules (Macías-Sánchez et al., 2017; Seto et al., 2012). These studies also showed that the darker phase, when preserved adequately, is distributed as more or less continuous pellicles. The thread-like distribution we observe is most likely due to deficient preservation of the observed growth surfaces. Both AFM and TEM observations are consistent with the presence of a minor ACC component.

**Evolution of the organo-mineral pattern during growth**

The study by focused ion beam coupled with SEM (FIB-SEM)-aided slice and view allowed us to track the evolution of the organo-mineral network during growth. By merely observing the successive slices, one can appreciate that the network is in a permanent dynamic state, even for reduced thicknesses (scanned volume ZEISScan 1; see method details). Polygons can undergo three different processes: (1) enlarging, (2) shrinking (to disappearance), and (3) division into two or more daughter polygonal units by intercalation of new membranes. Particular examples can be seen in Figures 4A–4C and Video S1. When a membrane emerges from one of the sides, the latter, being initially flat, bends at the junction, such that the initial 90°-90°-180° junction tends to transform progressively into a triple junction at 120° (Figure 4C). The quadruple points observed are not permanent and split with growth into two separate triple points (Figure 4D). All these processes do not take place immediately, but after a certain, sometimes prolonged, growth (i.e., thickness) delay.

In order to determine if these processes obey particular rules, we tracked the changes in surface area of individual polygons within two selected groups (seven polygons initially each) during the growth of the fibrous layers in each of the scans TESScan 1 and ZEISScan 1 (see method details). The curves obtained show that small polygons (below a cross-sectional area threshold value) tend to shrink until disappearance, whereas large polygons tend to become larger (Figures 5A and S2). The threshold in the four groups selected is 0.2–0.3 μm² (Figures 5 and S2). Of interest, particularly large polygons tend to split due to the appearance of new membranes. In this way, the polygons split into two or, more rarely, three, sometimes in a repeated fashion (Figures 5B and S2A). In general, there is not a clear-cut threshold value for polygon splitting: the minimum values observed are in the order of 0.5 μm² (Figure S2B), although other polygons do not split until values ≥2 μm² (Figures 5A and S2).

The group of polygons shown in Figure S2B was selected close to a neighbor spherulite. Owing to the cut being oblique, the neighbor progressively invaded the group and gradually outcompeted the polygons. Some polygons became temporarily wholly incorporated within the neighbor before disappearance (Figure 4E, and two right crops at the top of Figure S2B). This explains why all ascending curves eventually acquired a negative slope, soon after they were reached by the polygons of the advancing front of the neighbor spherulite (blue arrows in Figure S2B).
For the whole ZEISScan1 area, containing hundreds of polygons, we have calculated the number of polygons (N), the mean surface area (Average Area) (Figure 6A), and the standard deviation (SD) (Figure 6B). Since SD is dependent on the mean size, we have also calculated the coefficient of variation (CV = SD/Average Area) (Figure 6B) as a measure of size distribution independent of the scale. A plot of these parameters versus the increase in thickness reveals that they all remain relatively constant during growth (Figure 6). If we replot these parameters after manually eliminating the newly formed membranes (i.e., by suppressing polygon splitting)

Figure 3. TEM and AFM analysis of the calcite of Argonauta hians
(A and B) TEM analysis of two particles obtained through grinding. The particles are the small upper insets and the areas analyzed are framed in red. In both cases, the particles are crystalline but the rims appear amorphous, as indicated by the high-resolution images and the fast Fourier transform (FFT) of the framed areas. In the particle in (B), the amorphous rim recrystallized (spots in the FFT) after 15 min of irradiation. 
(C and D) AFM images of two different areas of the growth surface at different magnifications. The surfaces of the prisms display a characteristic roughness. The higher-magnification views in (D) show the existence of a low contrast (yellow-orange) background phase and a high-contrast (dark brown) phase, in the form of threads (arrows in phase image). The amplitude image shows that the threads are distributed between the nanoelevations of the surface (arrows in amplitude image).
In each case, a 3D view is provided, together with three sections at different depths (bottom, top, and intermediate; red arrows)
(A) Example of polygon enlarging and shrinking. The two lateral polygons enlarge, while the central element (the smallest one) shrinks until disappearance.
(B and C) Examples of polygon splitting into three (B) or four (C). The thick arrows in (C) indicate transformation of the initially flat side into a triple joint due to the appearance of a new membrane. Growth direction is toward the background in the two cases.
(D) Transformation of a quadruple point into two triple points.
(E) Detachment of two polygons (1 and 2) from a spherulite and incorporation within the neighbor spherulite; gf is the growth front of the spherulite to which the two elements belong. Polygons 1 and 2 are the same as 7 and 4a in Figure S2B, respectively. Long white arrows in (A, D, and E) indicate the growth directions. Growth in (B and C) is toward the background. (A–D) were obtained from the scanned volume ZEISScan 1, whereas (E) was obtained from TESScan 1. The volume represented corresponds to the mineral infillings in (A) and to the organic sheaths in (B–E).

(eliminated membranes can be seen in Figure S3), there is a downward trend for N and a drastic upward trend for Average Area (curves labeled “nmr” in Figure 6), as expected. At the same time, SD increases and CV decreases slightly. This is easily explained because, in normal conditions, the splitting of large polygons compensates for the disappearance of small polygons, thus providing constant N and Average Area and keeping SD more or less constant. In the absence of splitting, large polygons continue to enlarge, thus accentuating the size differences (i.e., higher SD). The slight decrease of CV in the absence of splitting is caused by the increase of Average Area being somewhat faster than that of SD. Our procedure of eliminating the new divisions through the different slices does not provide a fully realistic solution, since splitting interrupts the trend of large polygons to continue growing. Splitting occasionally leads to particularly small polygons, which shrink with time (e.g., polygonal element 2b in Figure 5A and polygonal elements 2ba and 2bbb in Figure 5B). Foreseeably, in the absence of splitting, Average Area and SD should increase above our calculations. We cannot quite predict the behavior of CV since it depends on the change speed of the above two parameters. In summary, the splitting process of large polygons is responsible for keeping all the measured parameters constant throughout growth. All parameters, except for CV, correlated significantly with the thickness (Table S2).

A similar calculation was carried out on the much larger and deeper TESScan 2 volume (containing thousands of polygons and a depth of ~32 µm). The plots obtained (Figure S4) display accentuated fluctuations in the above parameters. This is due to the slicing going through growth bands with particularly small polygons, i.e., at a larger scale there are growth rhythms marked by changes in the relative productions of the organic/mineral phases.

**Crystallography and distribution of the calcite phase**

Electron backscatter diffraction (EBSD) orientation maps indicate that the calcite crystals of both the outer and inner fibrous layers are broadly oriented with their c-axes perpendicular to the outer and inner surfaces,
Figure 5. Surface area versus thickness plots of selected groups of polygonal elements (PEs) (A and B) were selected from the TESScan 1 and ZEISScan 1 sliced volumes, respectively. The gray bands indicate the approximate size boundary above or below which PEs shrink or enlarge, respectively. The pink bands mark the lowermost and uppermost cases of splitting. Splitting events are indicated with thick arrows. Note splitting into three of PE 2a (arrow labeled with 3) in (A), and repeated splitting of PE 2 and its daughter PEs in (B). The upper crops display the distribution of PEs at different thicknesses. See additional examples in Figure S2.
i.e., parallel to the growth direction (Figure 7A, left panel). The a-axes can take on any orientation around the c-axis. The resulting texture is a fiber or axial texture. Each spherulite initially consists of a few crystals whose c-axes are arranged in a fan-like orientation (Figure 7A, right panel). With growth, the crystals situated more to the sides of the spherulites disappear due to the impingement onto the neighbor spherulites. In this way, both the number of crystals composing a given spherulite and the spread of their c-axes become reduced with growth. All in all, there is a greater spread of orientations close to the central organic layer, which soon becomes reduced with the impingement onto adjacent spherulites.

The cropping of particular domains, each identifiable by a reduced three-dimensional crystallographic orientation spread (see method details), reveals their complex outlines. In sections transverse to the growth direction (Figure 7A, left panel). The a-axes can take on any orientation around the c-axis. The resulting texture is a fiber or axial texture. Each spherulite initially consists of a few crystals whose c-axes are arranged in a fan-like orientation (Figure 7A, right panel). With growth, the crystals situated more to the sides of the spherulites disappear due to the impingement onto the neighbor spherulites. In this way, both the number of crystals composing a given spherulite and the spread of their c-axes become reduced with growth. All in all, there is a greater spread of orientations close to the central organic layer, which soon becomes reduced with the impingement onto adjacent spherulites.

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Figure 7. Crystallography of the calcite phase and its spatial relation with the organic framework

The crystallographic indices ((001) for calcite c-axes and (100) for calcite a*-axes) are Miller indices (i.e., plane normals) referring to the "hexagonal" unit cell of calcite. The color images are representations of crystallographic orientations, the latter are shown color coded, relative to a defined color code (color triangles in A and B).

(A) EBSD map of a section perpendicular to the shell surface across the middle organic layer (left). The pole figures indicate that the c-axes are parallel to the axes of elongation of the fibers. In the right panel, five spherulites have been cropped out and analyzed individually. Analysis of the lattice orientation within domains indicates that the c-axes within each spherulite have a radial orientation. The spreads of the c-axis orientation are indicated in the (001) pole figures with red arrows.

(B) EBSD map of a section transversal to the fibers. Some domains (each characterized by a particular crystallographic orientation) have been isolated. They have very complex, fractal-like outlines. Some of them have engulfed portions of other domains (white arrows).
The crystals presumably nucleate onto the organic center and must already be bounded by the formed organic membranes at this early stage. We do not know if there is a one to one crystal domain/organic prism relationship. During further growth, some polygons and their domains will be eliminated, while the process of splitting will allow a given domain to extend over two or more polygons. Repeated splitting could explain why crystal domains extend over many polygons.

Crystal competition has been reported in biomineral aggregates (Crippa et al., 2020; Goetz et al., 2009; Rodriguez-Navarro and Garcia-Ruiz, 2000) and was invoked also in the shell of Argonauta hians based on an increase of the EBSD-determined multiple of uniform distribution (MUD) values (which are a measure of crystal co-orientation; see method details) with growth (e.g., Stevens et al., 2017). Nevertheless, direct crystal contact (i.e., competition) is not expected since the calcite crystals are confined within the organic membranes. In fact, the outlines of crystal domains are not of the crystalline type (Figures 7 and S5). An increase in the MUD values can also be expected from the competition observed at the boundaries between spherulites, where the more oblique fibers containing the more inclined crystals (Figure 7A, right panel) are progressively eliminated with growth (Figures 1B, 1D, and 1E), i.e., competition between polygons may lead to an apparent competition between crystalline domains.

DISCUSSION

The shell mineralogy of the female Argonauta is fully calcitic, with a high amount of Mg (<10% molar substitution). This is 2-fold the previously reported value (Wolfe et al., 2012, 2013), and several-fold above any other molluscan calcite, always low-Mg calcite (e.g., Freitas et al., 2005). The amount of organic matter calculated by TGA-DSC (<8.5%) is notably higher than the amount calculated by XRD, and is in striking contrast with the ~1.8% of organic matter calculated by Oudot et al. (2020) after extraction by dissolution and ultrafiltration. The latter might be a low estimate since part of the biomolecules may have been lost during extraction. FTIR data indicate a significant predominance of proteins and is in agreement with previous results on the acid-insoluble matrix fraction of Argonauta hians (Oudot et al., 2020).

The shell of the female Argonauta is not secreted by the mantle of the animal, as in all mollusks, but by a completely separate tissue: the webs of the two dorsal arms. Accordingly, it is not homologous to the shell of mollusks, but an innovation of the Argonautidae, dating from the Late Oligocene (Noda et al., 1986). Hence, it is not surprising that, compared with the shells of mollusks, it is unusual in (1) its fully calcitic nature (calcitic bivalves, such as oysters, retain an aragonitic myostracum and intraligamentary fibers), (2) the high-Mg chemistry of the calcite, and (3) the bidirectional growth (toward the interior and the exterior of the shell) of the fibrous layers.

The organic network, when observed in transversal section, constitutes a polygonal cellular pattern with a minimal energy configuration, characterized by triple junctions at ~120° (Plateau’s law). We have also found that the polygonal network follows the von Neumann-Mullins topological law (Mullins, 1956; Von Neumann, 1952) during growth, since the individual polygons above a certain size (cross-sectional area) threshold (estimated here at 0.2–0.3 μm²) continue to enlarge, whereas those below that threshold shrink until disappearance. Similar behavior was demonstrated in the calcitic prismatic shell of the bivalve Pinna nobilis (Bayerlein et al., 2014), which was interpreted as a case of normal grain growth. Alternatively, Checa et al. (2016a) attributed such behavior in this and related species to the thick organic membranes surrounding the mineral units. They proposed that the system evolves as a biliquid foam or emulsion created between the fluid precursors of both the mineral, the so-called PILP (Olszta et al., 2003; Schenk et al., 2012), and the organic phases (gel-like precursor). In the prismatic layers of bivalves, given the relative
dimensions of the extrapallial space (EPS) (~100 nm thick), the mineral prismatic units (tens of micrometers in diameter), and the organic membranes (1–5 μm in thickness), it seemed unlikely that the mineral and organic components, if poured together to the EPS, could diffuse to their corresponding positions in the mineral units and organic membranes, respectively. Also, given the diameters (<10 μm) of the outer mantle cells (OMCs), below the mineral units’ diameters and above the organic membranes’ thicknesses, several OMCs must cooperate in the secretion of a single prismatic unit. Some OMCs would fall entirely below a mineral unit and would only secrete mineral components. Conversely, others would be in contact with the intervening organic membrane and would secrete either mineral or organic precursors in different areas of their surface, depending on which material they are in contact with. Accordingly, OMCs must recognize the substrate previously secreted, through contact recognition, and use it as a template for further secretion. This kind of secretion requires strict orchestration of the adjoining OMCs. Similar cellular recognition processes were invoked for the formation of the calcitic fibrous microstructure of brachiopods (Simonet Roda et al., 2019), the aragonitic helical fibrous microstructure of some planktonic gastropods (Checa et al., 2016b), and the vesicular microstructure of some oysters (Checa et al., 2020). Substrate recognition by cells is widespread in biological processes, like cell aggregation, growth, or immune response (Culp, 1978; Obrink and Ocklind, 1983; Saier and Jacobson, 1984). Recognition is carried out by cell surface receptors, able to identify ligands (both receptors and ligands are particular biomolecules) present in the substrate. This mechanism is also developed by mollusks (Gerdol et al., 2018). It is exploited in biomimetic science to develop bioadhesive and biocompatible materials by coating their surfaces with cell-recognizable natural proteins (Rahmany and Van Dyke, 2013).

We have demonstrated that in the shell of Argonauta the organic framework dominates the pattern and evolves following the laws of emulsions, foams, and metallic grain aggregates. As explained above for the prismatic layers of bivalves (Checa et al., 2016a), of these three possibilities only a system fluid-fluid is biophysically possible, that is an emulsion between the mineral and organic fluid precursors created at the interface between the mineralized shell and the living tissue (Figure 8). Our ultrastructural AFM and TEM data on the calcite of Argonauta indicate that its nanostructure is identical to that of other biominerals, including the presence of residual ACC. Accordingly, we can safely assume that the high-Mg calcite crystallizes from an ACC precursor, as in invertebrate biocrystals (Addadi et al., 2003), most likely in the form of a PILP, generated by acidic biopolymers (Olszta et al., 2003). It has recently been found that the PILP for calcium carbonate consists of ACC nanoclusters, with liquid-like behavior (Xu et al., 2018), i.e., it is a colloidal phase. The organic phase (the membrane network) would constitute the continuous phase of the emulsion. The membranes are constituted by highly organized self-assembled double layers of fibrous proteins. Proteins play important roles in emulsions, particularly as stabilizers (e.g., in Pickering-type emulsions) (Cabra et al., 2008; Hoffman and Roë, 2014), to which they impart gel-like properties (emulgels) (Glusac et al., 2018; Sun and Holley, 2011).

In Argonauta, virtually nothing is known about the characteristics of the biomineralization system created between the forming shell and the webs of the dorsal arms. Nevertheless, given the smoothness and continuity of the growth lines (Figures 1A, 1B, and 1D–1F), we may hypothesize that the dorsal webs must be very close to the shell growth surface during secretion (Figure 8), in a way similar to the mantle of the rest of shell-forming mollusks. Compared with the prismatic layers of bivalves, the surface areas of polygons in Argonauta (rarely above 1 μm²) are much smaller than those of the biomineralizing living cells of the dorsal webs (DWCs). Although there may be some doubt on the involvement of contact recognition processes in the formation of the argonaut shell, there is a process unique to it that makes such processes ineluctable: polygon splitting. This feature cannot be driven physically because it acts against the reduction of the free energy of the system (the number of interfaces), according to the von Neumann-Mullins law. Polygon splitting undeniably demands that the living DWCs are able to locate the polygons exceeding a certain cross-sectional area and extend new membranes across them (Figure 8). Note that the membranes in Argonauta are much thinner (>100 nm) than those of the prismatic layers of bivalves (1 to several micrometers) (Checa et al., 2016a), which speaks about the sensitive abilities of the DWCs, i.e., recognition and secretion takes place well at the submicron level. Given the continuity of the organic framework, there must be a coordinated and refined recognition-secretion activity of the mineralizing DWCs. Frequently, the production of new membranes and the detection of large polygons will be shared by two or more neighbor DWCs, since those objects do not necessarily fall within the ambitus of single DWCs. Accordingly, the implicated DWCs must act in coordination to carry out their secretory/sensitive activity. A recent proteomic study (Oudot et al., 2020) found that the organic matrix of Argonauta hians contained several hits with proteins that bind to signaling molecules. These might easily derive from the DWC activity we invoke here.
We found that the threshold for enlargement/shrinking of the polygons is narrow (calculated here between 0.2 and 0.3 m²), whereas that for polygon partition is very wide (0.7–2.2 m²). The difference lies in that the former is determined by a physical process (emulsion coarsening), whereas the second has a biological basis (location of particularly large polygons by the DWCs and subsequent secretion), with a much wider control range.

Polygon splitting clearly aims at maintaining mean cross-sectional surface area (Average Area) and area distribution (SD and CV; the latter being called polydispersity in colloidal systems) constant throughout growth. There might be a functional reason behind it. The organic membranes of the prismatic layers of bivalves make the material highly elastic. These materials constitute the margins of bivalves of the order Ostreida and flex upon valve closure, thus providing a tight marginal seal (Harper and Checa, 2020). Power (1856) reported that the shell of Argonauta argo is also highly flexible. Accordingly, maintaining the density of organic interfaces constant (by polygon splitting) would permit to retain the flexibility upon shell thickening.

Limitations of the study

Technically, the study was limited by the quality of the FIB-prepared surfaces. Despite the use of a rocking stage to minimize the curtain effect (see method details), some of the images had to be manipulated by hand, owing to the incompleteness of the organic boundaries. From the conceptual viewpoint, a main shortcoming is the virtual absence of information about the biominalization system in Argonauta, particularly with regard to the structure of the biominalizing tissue of the dorsal webs and its position with respect to the shell surfaces. We had neither access to living specimens nor the means to keep them in aquarium. Even if this had been the case, we would have been faced with the almost impossibility to keep the webs in position during specimen preparation for histological study. The inference we make of an extremely thin extrapallial space is based on the comparison with some known cases of the molluscan mantle, and on the evenness of growth surfaces. There is no expectation that this gap in knowledge can be filled in the short to medium term.

Figure 8. Evolution of the organo-mineral pattern during growth of the fibrous layer of Argonauta hians and model for its fabrication

For a Figure360 author presentation of this figure, see https://doi.org/10.1016/j.isci.2021.103288.
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SUPPLEMENTAL INFORMATION
Supplemental information can be found online at https://doi.org/10.1016/j.isci.2021.103288.

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AUTHOR CONTRIBUTIONS

DECLARATION OF INTERESTS
The authors declare no competing interests.

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REFERENCES


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KEY RESOURCES TABLE

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RESOURCE AVAILABILITY

Lead contact
Further information and requests for data should be directed to and will be provided by the Lead Contact, Antonio G. Checa (acheca@ugr.es).

Materials availability
This study did not generate new reagents or materials.

Data and code availability
The data can be obtained from the Lead Contact, Antonio G. Checa (acheca@ugr.es), or the coauthor Fátima Linares (flinaor@ugr.es).

Raw quantitative area measurements from Figures 5, 6, and S2, and raw crystallographic (EBSD) data (*.cpr and *.crc files generated by AZtec software) from Figures 7 and S5 were deposited in Mendeley Data: https://data.mendeley.com/datasets/n37bddwy7k/3.

Any additional information required to reanalyze the data reported in this work paper is available from the Lead Contact upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Shells
Four empty shells of Argonauta hians were purchased from Conchology Inc. (https://www.conchology.be/), two of which were used for analysis. They all came from the Philippines (exact localities unknown). They are not protected by the Convention on International Trade in Endangered Species of Wild Fauna and Flora (CITES).
METHOD DETAILS

X-Ray diffraction (XRD)
Shell fragments were gently ground to a fine powder with an agate mortar and pestle and mixed with silicon powder (42.26 wt%). The mineralogy of the shell was analyzed with a Panalytical Xpert Pro X-ray powder diffractometer from the Department of Mineralogy of the University of Granada (UGR) (Cu Kα; 2θ range: 4° to 120°; 0.013° step size; 100 s per step). To study the contribution of shell organic matrix to calcite lattice, the same sample was analyzed again after heating at 400°C and annealed for 30 mins. Calcite unit cell parameters and the percentage of amorphous were determined by Rietveld refinement using TOPAS 5.0 software (Bruker, Germany), considering calcite and silicon as the internal standard.

Thermogravimetry (TGA) and differential scanning calorimetry (DSC)
About 25 mg of powdered shell material were analyzed with a TGA and DSC coupled system from METTLER-TOLEDO (model TGA/DSC1) housed at the Center for Scientific Instrumentation (CIC) of the UGR. A heating rate of 10°C/min in air or nitrogen was used for registering the TGA/DSC scans.

Fourier-transform infrared spectroscopy (FTIR)
One shell fragment (~1 cm²) was immersed in a decalciying and etching solution (0.05M EDTA + 0.25 M HEPES + 2.5% Glutaraldehyde) until complete decalcification. The organic residue was washed with ultrapure distilled water (milli Q) and oven-dried. It was analyzed with an attenuated total reflection (ATR) diamond unit (ATR Pro One, Jasco) (Department of Mineralogy, UGR). The IR spectra were recorded at a 2 cm⁻¹ resolution over 100 scans using a Fourier transform infrared (FTIR) spectrometer Jasco 6600. Four measurements were carried out in different areas of the membrane to obtain a coherent assertion (standard deviation and error) of the results.

Atomic force microscopy (AFM)
Analyses were done directly on both the outer and inner surfaces of shell fragments, previously cleaned with sodium laurel sulphate. We used an AFM Park Systems NX20 (CIC, UGR) equipped with a cantilever ACTA (K= 40 N/m, F= 280 kHz) (CIC, UGR). Scans were done in tapping mode while displaying height, amplitude and phase signals. Images were obtained with Smart Scan v12 and subsequently analyzed with XEI 4.3 software (Park Systems).

Scanning electron microscopy (SEM)
Ultrasonicated fragments were cleaned by immersion in commercial bleach (~5% active chlorine) for 10 min. Successive washings in MilliQ ultrapure water, for 2-3 min each, followed. Dried fragments were embedded in epoxy resin (EpoFix, Struers), sanded, and polished in successive steps until a mirror-like polish was obtained with 0.25 μm diamond suspension (Struers). Some of the polished samples were etched (2.5% glutaraldehyde, 0.25M HEPES buffer, and 0.05M EDTA), for 2 min in an orbital shaker. Other samples were exposed to the etching/fixing solution for much longer times to completely denudate the organic network. All samples were carbon-coated (Emitech K975X carbon evaporator) and observed in the field emission SEMs (FESEM) Zeiss Auriga and FEI QemScan 650 F (CIC, UGR).

Electron backscatter diffraction (EBSD)
Shell samples were embedded in epoxy resin and subjected to several sequential grinding and polishing steps. The two final steps consisted of etch-polishing with colloidal alumina in a vibratory polisher and subsequent polishing with an Ar beam in an ion polisher. The samples were coated with 4-6 nm of carbon. Measurements were carried out on a Hitachi SU5000 field emission SEM, equipped with an Oxford EBSD detector (DEES, LMU). The SEM was operated at 20 kV. Data acquisition (with step sizes of 150 and 300 nm) was done with the Aztec (Oxford Instruments) software.

Information obtained from EBSD measurements is presented as grey-scaled band contrast measurement images and as color-coded crystal orientation maps with corresponding pole figures; the latter giving either individual data points or contours. In the latter case, we use the lowest possible degree for half-width (5°) and cluster size (3°). The half-width controls the extent of the spread of the poles over the surface of the project sphere. A cluster (or domain) comprises data with the same orientation. The color code used for crystal orientation is either given in the figure or stated in the figure caption. EBSD band contrast gives the signal strength of the EBSD-Kikuchi diffraction pattern at each measurement point and is shown as a
grey-scale component in a map. When the strength of the EBSD signal is high, we find bright gray colors in the map, whereas when it is weak or absent, gray colors are dark. Crystal co-orientation statistics are derived from density distributions in pole figures and are given as MUD (multiple of uniform distribution) values. The higher the MUD, the higher the co-orientation. All post-processing was done with the Oxford Instruments CHANNEL 5 HKL software.

3D reconstruction by focused ion beam coupled to SEM (FIB-SEM)

Three volumes were reconstructed by using the slice and view technique by means of FIB-SEM. Two of them were scanned with a TESCAN Amber X FESEM equipped with Plasma FIB (TESCAN Central Demo Lab & Applications, Brno, Czech Republic). Polishing was done using a current of 10 nA, and a rocking stage was used to reduce the curtaining effect. Images were acquired in backscatter electron (BSE) mode at 5 kV. The first scanned volume (TESScan 1), with dimensions (width x height x depth) 25x30x29 μm, consisted of 182 slices with a thickness of 50 nm, oblique to the growth direction of the fibers. The second volume (TESScan 2) (50.5x32.4x31.9 μm) consisted of 525 slices perpendicular to the growth direction; slice thickness was 60 nm. A third volume (ZEISScan 1) was analyzed with a Zeiss Crossbeam 350 FESEM, equipped with a gallium FIB column (Carl Zeiss Microscopy GmbH, Oberkochen, Germany). The polishing current was 50 pA. Images were collected in Energy Selective Backscattered (EsB) mode. The volume scanned (12x12x3.6 μm) generated 360 images perpendicular to the growth direction, separated by a thickness of 10 nm.

Area analyses were done with FIJI software (Image J). Raw TIFF images were converted to 8-bit images. After B/W threshold adjustment, we applied Noise-Despeckle followed by Binary-Watershed. Before particle analysis, polygon boundaries of TESScan 1 and ZEISScan 1 images were manually retouched for maximal accuracy. Given the high number of polygons (thousands) contained within the images of TESScan 2, we skipped this procedure. Video, segmentations and 3D meshes were done with Dragonfly™ 4.1 software after slice registration (registration method SSD). All the procedures were carried out at the CIC (UGR).

Transmission electron microscopy (TEM)

The surface of a shell fragment was scraped with a scalpel. Fragments were ground in an agate mortar and the powder was suspended and sonicated in pure ethanol. The solution was dropped onto lacy carbon copper grids and air-dried for subsequent TEM observation. Measurements were acquired using a double Cs corrected FEI Titan G2 60-300 TEM equipped with an X-field emission gun and a Gatan Ultrascan camera (CIC, UGR). Imaging was performed in TEM mode at 300 kV, and 0.5-1 s exposure. Nanodomains were indexed by Fast Fourier Transform (FFT) analysis.

Other shell fragments, were immersed in a fixing decalcifying solution (0.05 M EDTA, 2.5% glutaraldehyde, and 0.25M HEPES) until complete decalcification occurred (48-72 h). The residual membranes were critical-point dried (Leica EM CPD300), post-fixed in OsO4 (2%) for 2 h at 4°C and embedded in epoxy resin Epon 812 (Electron Microscopy Science, EMS). Ultra-thin sections (50 nm), obtained with an ultramicrotome Leica Ultracut R, were stained with uranyl acetate (1%), followed by lead citrate. They were later carbon-coated and observed with TEM Zeiss Libra 120Plus (CIC, UGR).

Illustration

All figures were illustrated by using the Corel Draw Graphics Suite (version Home & Student X8).

Video production

Video production for TESScan 1, TESScan 2, and ZEISScan 1 datasets (see subsection “3D reconstruction by focused ion beam coupled to SEM (FIB-SEM)” above for details) was performed with Dragonfly 4.1 Movie Maker. Only the video generated from ZEISScan 1 dataset is presented, as Video S1.

QUANTIFICATION AND STATISTICAL ANALYSIS

Statistical analyses were made with Statistical Table Calculator (http://vassarstats.net/tabs_r.html).