



Protective role of *Arapaima gigas* fish scales: Structure and mechanical behavior



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ABSTRACT

The scales of the arapaima (*Arapaima gigas*), one of the largest freshwater fish in the world, can serve as inspiration for the design of flexible dermal armor. Each scale is composed of two layers: a laminate composite of parallel collagen fibrils and a hard, highly mineralized surface layer. We review the structure of the arapaima scales and examine the functions of the different layers, focusing on the mechanical behavior, including tension and penetration of the scales, with and without the highly mineralized outer layer. We show that the fracture of the mineral and the stretching, rotation and delamination of collagen fibrils dissipate a significant amount of energy prior to catastrophic failure, providing high toughness and resistance to penetration by predator teeth. We show that the arapaima's scale has evolved to minimize damage from penetration by predator teeth through a Bouligand-like arrangement of successive layers, each consisting of parallel collagen fibrils with different orientations. This inhibits crack propagation and restricts damage to an area adjoining the penetration. The flexibility of the lamellae is instrumental to the redistribution of the compressive stresses in the underlying tissue, decreasing the severity of the concentrated load produced by the action of a tooth. The experimental results, combined with small-angle X-ray scattering characterization and molecular dynamics simulations, provide a complete picture of the mechanisms of deformation, delamination and rotation of the lamellae during tensile extension of the scale.

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1. Introduction

Natural protective layers, exhibited by seashells, fish scales, turtle carapaces, armadillo and alligator osteoderms, have developed and refined over hundreds of millions of years through a process of convergent evolution [1]. These biological materials have hierarchical architectures and exhibit excellent properties with complex mechanisms of failure avoidance. They are inspiring man-made

structural materials that are lightweight and demonstrate outstanding toughness; good examples are nacre-inspired, freeze-cast structures [2–4].

Dermal protection of animals ranges from rigid to flexible. Flexibility enables mobility while rigid components maximize protection. Fish scales uniquely combine these two traits, and can be classified into four groups: placoid [5,6], ganoid [7–9], cosmoid [10–12] and elasmoid (cycloid and ctenoid) [13–16]. Yang et al. [1] reviewed the four types and provided a schematic drawing of their morphologies as well as their connection mechanisms. Placoid scales are denticles, or small modified teeth, which cover the cartilaginous fish's skin. They give the skin a rough feel, and may contribute to favorable water flow while swimming. Ganoid scales are recognized by a thick surface layer of enamel or ganoine, on top

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of a dentin or bony base. Cosmoid scales consist of a double bone layer, comprising lamellar and vascular bone; the outer layer is considered as a dentin-like cosmine. Elasmoid scales are thin, lamellar collagenous plates. Cycloid scales are composed of concentric rings and ctenoid scales are composed of fringed projections along the posterior edge. In fish, scales are lightweight and enable flexibility to swim, yet are sufficiently rigid to provide protection from predators.

Most fish scales have similar constituent materials: calcium-deficient hydroxyapatite (HAP) and type I collagen fibrils [17–19]. The mineralized collagen fibrils form different architectures, such as the plywood (Bouligand-type) structure [20–22] or, in the case of ganoid scales, a bony structure with a complex structure interlaced with HAP crystals [7,8]. Additionally, scales commonly exhibit a composite or graded structure with hardness decreasing from the outer to the inner layers.

Studies on the penetration of scales using either real teeth or simulated indenters to mimic fish bites provide a good estimate of the effectiveness of the scale's complex structure. Zhu et al. [16] penetrated single scales of striped bass (*Morone saxatilis*) with a sharp indenter, and analyzed the sequence of events, dividing it into three stages: elastic deflection of the scale, fracture of the mineralized layer and penetration of the collagen lamellae. Meyers et al. [23] penetrated the arapaima scale with a real piranha tooth since piranha is the major predator to arapaima in the Amazon River; despite being twice the hardness of the scales, the tooth failed to penetrate them. Zhu et al. [24] found that overlapping three scales essentially multiplies the puncture resistance by three. Friction between the scales is negligible, and therefore does not generate additional resistance to deformation or puncture, regardless of the arrangement of the scales. The force dispersal from scales prevents unstable localized deformation of the skin and damage to underlying tissues. Finite element modeling of the penetration into ganoid [25] and elasmoid [26] scales was conducted to establish the interactions between tooth sharpness, scale flexibility and other parameters.

It was established by Lin et al. [22], Meyers et al. [23] and Zimmermann et al. [27] that the structure of the arapaima scale consists of an external layer with a highly mineralized, rough surface and inner foundation of collagen layers organized in a Bouligand-type structure with lamellae (~50–60 μm thick for each) of parallel collagen fibrils in different orientations. However, the understanding of the fundamental mechanisms of deformation and damage avoidance in elasmoid scales is still incomplete. The objective of this work is to present a comprehensive evaluation of the mechanisms governing the strength, ductility and toughness of the scales of the arapaima fish (Fig. 1a) through three complementary methods: experimental, analytical and computational. The damage mechanisms of the penetration by teeth are identified and quantified through controlled experiments. The principal experimental method is the *in situ* observation of the deformation of a scale. This is supplemented by an analysis based on the interaction between the different lamellae and by molecular dynamics (MD) calculations.

2. Methods

The arapaima scales (Fig. 1b), which typically are 50–120 mm in size, were characterized using a range of experimental procedures to determine their structure and mechanical performance. In particular, indentation and tensile tests were performed on whole scales and on the collagen layer alone (after removing the outer mineral layer) to establish the individual function as well as deformation and fracture mechanisms in each layer. Additionally, the local behavior of the collagen was investigated using *in situ*

small-angle X-ray scattering (SAXS) studies during uniaxial tensile testing; this behavior was also analyzed using MD techniques to quantitatively predict the controlling damage mechanisms in the inner layer.

2.1. Characterization of scales

The structure of the scales, as well as failure mechanisms and distribution of damage after mechanical testing, were investigated in an FEI SFEG ultrahigh-resolution scanning electron microscope (SEM; FEI, Hillsboro, OR), Veeco scanning probe microscope (Veeco, Plainview, NY) and an FEI Tecnai 12 transmission electron microscope (TEM) operating at 120 kV. All samples for SEM characterization were sputtered with iridium prior to observation.

2.2. Penetration and indentation

The penetration resistance of the entire scale including the external layer was examined using an Instron 3367 mechanical testing machine (Instron Corp., Norwich, MA) with a load cell of 30 kN. The scales in fish are in general designed to resist penetration by teeth. In the case of the arapaima scales, this necessity is extreme due to the sharp teeth of their principal predator, the piranha. It was demonstrated by Meyers et al. [23] that the piranha tooth is unable to penetrate through the scales without being arrested or broken by the highly mineralized external layer of the scale. A piranha tooth was removed from the fish's head and attached to the upper punch in order to indent the entire arapaima scale (through the external layer), which was glued on a synthetic rubber layer with a compressive elastic modulus of ~10 MPa attached to the lower punch. The low modulus of the rubber simulates the flesh under the scale, which was established by Meyers et al. [23]. The penetration tests were performed at a displacement rate of 10 mm min^{-1} . Indentation of the inner layer (lamellar structure without mineralized surface) was carried out using an indentation load of 1 kgf and a holding time of 10 s using a LECO M-400-H1 (LECO, Joseph, MI) hardness testing machine. Both types of penetration/indentation tests were performed on wet scales (kept in water at 25 °C for 12 h).

2.3. Tensile behavior

Using a surgical blade, 15 × 2.3 mm tensile samples were cut in two orientations (Fig. 1) from the scales with thickness of 0.6–1.2 mm. The outer layers of some samples were removed using silicon carbide polishing paper of 180#–2500#, leading to final thickness between 0.6 and 0.8 mm (Fig. 1b). In order to prevent slippage, the ends of the samples were glued between sand paper sheets using cyanoacrylate glue, resulting in a gauge length of ~8 mm. Uniaxial tensile tests were carried out on an Instron 3342 mechanical testing machine (Instron Corp., Norwich, MA) with a load cell of 500 N at a strain rate of 10⁻² s⁻¹ immediately following the removal of samples from fresh water, where they were kept prior to testing. The tensile results were expressed in terms of engineering stress–strain curves. The ultimate tensile stress, uniform elongation (strain at maximum stress) and the toughness, as measured by the area beneath the stress–strain curves, were established.

2.4. *In situ* observation under tensile loading

To discern the salient energy absorbing mechanisms during penetration and to visualize the distribution of damage and corresponding resistance to cracking, both rectangular tensile samples and single-edge notched tension (SE(T)) samples, with and without the mineral layer, were prepared in the longitudinal direction with

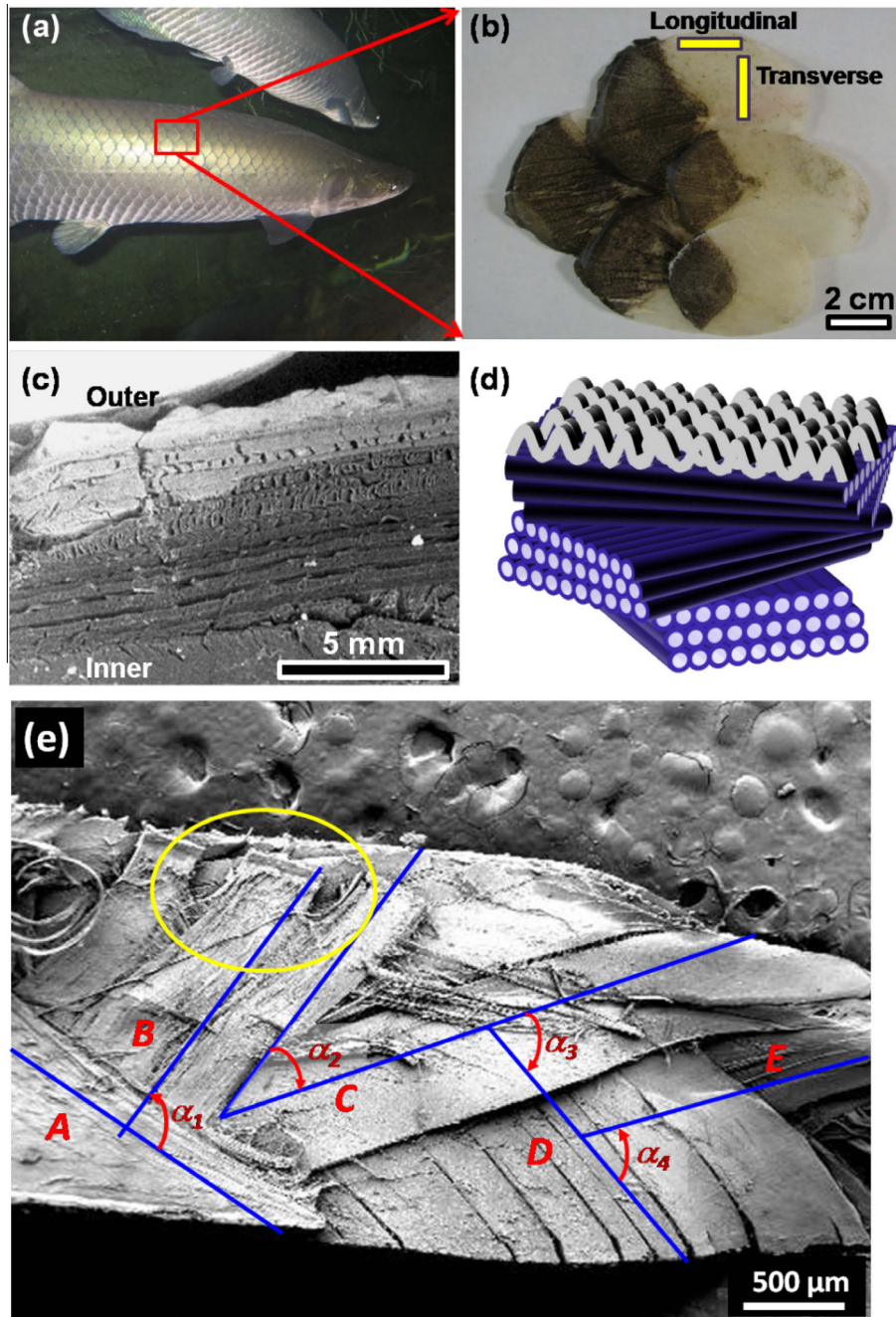


Fig. 1. Arapaima fish and its scales: (a) arapaima fish with overlapped scales on its body; (b) close-up of the overlapped scales (darker regions were exposed, lighter regions were covered by other scales) showing longitudinal and transverse directions of specimens for tensile tests; (c) scanning electron micrograph of the cross-section of a scale (note that the small cracks in the collagen layer are artifacts, formed when the scales dried out in vacuo); (d) schematic drawing of the structure of the arapaima's scale: the outer layer is highly mineralized, whereas the inner layer has different orientations of collagen fibrils arranged in lamellae; (e) tensile failure region of the lamellae: $\alpha_1 = 86.7^\circ$, $\alpha_2 = -34.5^\circ$, $\alpha_3 = -69.2^\circ$, $\alpha_4 = 66.9^\circ$. The yellow ellipse in (e) shows two events where lamellae separate leading to changes in the angle.

an 8 mm gauge length using the same dimension and procedure as in Section 2.3. The notch in the SE(T) samples was first cut using a diamond blade and sharpened by polishing with a razor blade irrigated in 1 μm water-based diamond suspension; the resulting micro-notch had a depth of approximately half the specimen width with a consistently sharp root radius of $\sim 5 \mu\text{m}$.

Uniaxial tension tests were performed at 25 $^\circ\text{C}$ in an environmental Hitachi S-4300SE/N (Hitachi America, Pleasanton, CA) SEM under wet conditions. To minimize any dehydration, the samples were soaked in water for at least 12 h prior to testing.

Testing was performed at a displacement rate of 0.5 mm min^{-1} using a Gatan Microtest 2 kN bending stage (Gatan, Abington, UK) inside the SEM; the samples were imaged in electron backscatter mode under variable low pressure conditions at a vacuum of 35 Pa. Such low pressure is often used for biological materials to limit the effect of dehydration and keep the properties of the material relatively unaltered; however, while the outer surface of the scale generally slightly dehydrates, the interior remains wet until exposed. Due to the excessive deformation during testing, as well as constant and significant relaxation during imaging,

a reliable measurement of load and displacement could not be obtained; consequently, the stress–strain curves, fracture toughness and crack-resistance curves (*R*-curves) could not be measured.

2.5. Small-angle X-ray scattering

Rectangular samples, 15 × 1 mm in size with ~0.6 mm thickness and gauge length of 10 mm, were sectioned with a razor blade using the same polishing procedure described above. The scales were hydrated in water for 12 h prior to mechanical testing.

The hydrated samples were loaded in uniaxial tension and exposed to X-rays at beamline 7.3.3 at the Advanced Light Source (ALS) synchrotron at the Lawrence Berkeley National Laboratory (Berkeley, CA). The mechanical tests were performed with a custom-made rig using a 10 mm displacement stage and an Omega LC703-10 load cell, calibrated to 45 N, to measure the force; this set-up permits SAXS data collection to be recorded in real time with the simultaneous measurement of the load–displacement curve. The mechanical tests were performed at 25 °C at a displacement rate of 1 μm s⁻¹, on hydrated samples maintained through the use of a hydration cell comprising a strip of cellophane held to the sample through capillary action of a few drops of water.

A Pilatus 1 M detector (Dectris Ltd, Baden, Switzerland), used to collect the SAXS data, was located at the largest allowable distance from the sample (~4 m) to permit detection of the fine changes in the collagen peak positions. The sample was exposed to X-rays for 0.5 s at ~5 s intervals during mechanical testing; exposures were in the less than 20 kGy range, below a level that would adversely degrade the tissue [28]. Further details concerning the data collection and analysis are given in Zimmermann et al. [27].

2.6. Preparation of TEM specimens

The preparation procedures used for biological specimens involved a number of stages. The arapaima scales were first immersion-fixed in 3% glutaraldehyde in 0.15 M sodium cacodylate buffer (pH 7.4) for 4 h, and then post-fixed using 1% OsO₄ solution with 8% potassium ferrocyanide in 0.15 M sodium cacodylate buffer for 12 h at 4 °C. The scales were subsequently stained with 2% aqueous uranyl acetate for 12 h and dehydrated with an ascending ethanol series (50, 70, 90 and 100%), followed by a 1/1 ratio of 100% ethanol and 100% acetone and finally 100% acetone. Samples were then embedded in Spurr's low viscosity resin and polymerized at 60 °C for 48 h. Samples were subsequently sectioned perpendicular to the scale surface to generate slices 80–200 nm thick using a Leica Ultracut UCT ultramicrotome (Leica, Wetzlar, Germany) and a Diatome diamond knife (Diatome, Hatfield, PA). The sections were post-stained with 1% uranyl acetate for 10 min and Sato lead for 1 min. At this stage they were ready for examination in an FEI Tecnai 12 TEM at 120 kV.

2.7. Preparation of atomic force microscope specimens

The lamellae with a thickness of ~200 μm were peeled off from the inner layer of arapaima fish scale. The inside peeled surfaces of the lamellae samples were probed in water with a Veeco scanning probe microscope using the SNL-10 Bruker tips (Bruker, Billerica, MA).

2.8. MD analysis

A MD model, based on a simple elastic network model [29,30], was used to theoretically investigate the mechanisms of the deformation and delamination in the arapaima scales under uniaxial tension. This model does not take the fibrils as elements but rather

considers the larger collagen fibers to better simulate the response of the lamellae. Based on Lin et al. [22], the initial inter-bead distance is $r_0 = 1 \mu\text{m}$, which is the same as the average diameter of the collagen fiber d_c . Each collagen fiber is modeled as a series of beads connected by harmonic springs. Since the density of the collagen is $1.34 \times 10^3 \text{ kg m}^{-3}$, the mass of each bead is equal to $1.05 \times 10^{-15} \text{ kg}$. The total deformation energy of the simulation system is given by:

$$U_{total} = U_T + U_B + U_{weak}. \quad (1)$$

As the total energy (U_{total}) is given by the sum over all pair wise (U_T), three-body (U_B) and weak inter-fiber interactions (U_{weak}) where $U_T = \sum_{\text{pair}} \varphi_T(r)$, $U_B = \sum_{\text{triplets}} \varphi_B(\theta)$ and $U_{weak} = \sum_{r < r_{\text{cutoff}}} \varphi_{weak}(r)$ with $\varphi_T(r)$ as the energy of inter-bead spring in tension, $\varphi_B(\theta)$ as the energy of angular spring in bending and interaction energy of inter-fiber beads. Here $\varphi_T(r) = K(r - r_0)^2$ is the energy of each inter-bead spring in tension, where K as the stiffness of the springs is assigned according to $K = \frac{EA}{2r_0}$ (estimated value of the Young's modulus of collagen fiber, $E = 2 \text{ GPa}$) and $A = \frac{\pi d_c^2}{4}$ is the cross-section area of the collagen fiber. An angular spring between two neighboring springs is used to define the bending stiffness of the collagen fiber, and its energy is given by $\varphi_B(\theta) = K_B(\theta - \pi)^2$, where the the bending stiffness of the thread is given by:

$$K_B = E I_t / (2r_0) = E \pi d_c^4 / (128r_0), \quad (2)$$

where I_t is the area moment of inertia of the cross-section. The inter-fiber interaction is denoted by:

$$\varphi_{weak}(r) = 4\epsilon \left[(\sigma/r)^{12} - (\sigma/r)^6 \right], \quad (3)$$

where ϵ is the interaction energy between two inter-fiber beads that is given by $\epsilon = d_c r_0 \gamma$ as γ is the surface energy of the collagen fibers, and $\sigma = d_c / \sqrt[5]{2}$ decides the equilibrium distance between two neighboring collagen fibers, which equals d for this selection. To ensure that the collagen fibers only interact with their nearest neighbors, the cutoff of the interaction is set as $r_{\text{cutoff}} = 1.5d_c$.

The model of the arapaima scale is composed of multiple layers of parallel collagen fibers [22] making specific angles which were found to vary from 35 to 85°. Considering that the mean angle between two neighboring lamellae is ~60°, it suffices to model three lamellae with 60° (one centrally sandwiched between two lamellae) to capture the most important events. For simplicity, each lamella is modeled by three layers of collagen fibers that initially align in the same direction; the thickness of each lamella is ~3 μm. This is much lower than the actual lamellar thickness, ~50 μm, but sufficient to capture the essential phenomena since we also include the periodic boundary conditions in the direction perpendicular to the lamellae.

We applied a uniaxial tensile strain to deform the entire multi-lamellar film in a quasi-static way. For every deformation increment, we applied a uniaxial strain of 0.004 to the model, fixed the single layer of beads at the two ends of the lamellae along the pulling direction and carried out energy minimization and equilibrium for 20,000 integration steps. We recorded the total force applied to the beads at the boundaries and obtained the stress–strain curve of the material during deformation. We also monitored the deformation of all the collagen fibers during the entire loading process. This model enabled a systematic investigation of how different mechanical and geometric characteristics of the arapaima scale influence its response to loading, e.g. the angle between two lamellae, the inter-fiber interaction strength and the stiffness of a single collagen fiber.

3. Results

3.1. Structure characterization

The cross-sectional structure of the entire arapaima scale is shown in Fig. 1c. Three parameters can be used to describe the geometrical characteristics of scales [31]: aspect ratio (= total length/thickness), degree of imbrication (= exposed length/total length) and the angle of scales with the surface plane. The (elasmoid) scales of this fish have an aspect ratio of ~ 50 and a degree of imbrication of 0.4 (Fig. 1a and b). They consist of hydroxyapatite and type I collagen arranged in two distinct layers: a rough, highly mineralized, outer layer and a poorly mineralized collagen inner layer (Fig. 1c). Our group [22] performed Fourier transform infrared spectroscopy, which showed several strong absorption peaks, similar to previous studies [14,32]. The absorption peaks at 1637, 1546 and 1239 cm^{-1} are the three characteristic peaks which correspond to amide I, amide II and amide III of type I collagen, respectively. Lin et al. [22] present a table comparing the peaks. There are also peaks at around 1000 cm^{-1} , which represent the phosphate groups at 872 cm^{-1} , peaks at 1401 and 1450 cm^{-1} correspond to carbonate anions. Within the inner layer, the fibrils align parallel within layers called lamellae that have a characteristic thickness of 50–60 μm [22,32]. Neighboring lamellae are rotated to form a twisted plywood structure similar to that proposed by Bouligand [33]; we refer to this as the Bouligand-type structure (Fig. 1d). Fig. 1e shows the extremity of a specimen that was stretched until complete failure (strain ~ 0.3); the arrangement of the different orientations of collagen fibrils in the Bouligand-type structure is clear. Collagen lamellae are labeled from the outside to the inside as A to E. It is apparent that the angles between the fibrils of adjacent lamellae are not equal, as would be the case for the classical Bouligand structure. Using the nomenclature defined in Fig. 1e, these angles are: $\alpha_1 = 86.7^\circ$, $\alpha_2 = -34.5^\circ$, $\alpha_3 = -69.2^\circ$, $\alpha_4 = 66.9^\circ$, i.e. displaying an irregular orientation which varies from lamella to lamella.

The properties of the scales are highly dependent on the level of hydration [22], due to the effect of water molecules on the bonding between fibrils, as computed by Gautieri et al. [34]. The calculated density of collagen decreases from 1.34 to 1.19 g cm^{-3} with hydration and is accompanied by a decrease in the Young's modulus from 3.26 to 0.6 GPa.

Fig. 2 shows the individual collagen fibrils imaged by TEM and atomic force microscopy (AFM). The fibrils imaged in Fig. 2a are not equiaxed because the foil plane is not exactly perpendicular to the fibril axis. The fibrils, initially cylindrical, become faceted because of the necessity of tight stacking and the lamellar orientation. In Fig. 2b two fibril orientations from two adjacent lamellae are imaged. The fibrils in the left image are close to perpendicular to the foil plane; the ones in the right image are close to parallel. The interface between lamellae with different orientations is abrupt, i.e. there is no gradual transition stage. The fibrils are cylindrical and arranged in the most compact fashion. The surfaces at which adjacent fibrils touch are somewhat flattened, enabling a tighter stacking. The apparent period in the collagen fibrils measured in the TEM micrograph of Fig. 2b is ~ 50 nm. The banding consists of three white radial lines, due to the fact that the foil thickness

(60–200 nm) can exceed the fibril diameter. Thus, two or three fibrils may simultaneously be imaged, creating the unusual pattern. The d period of the collagen in the right side of Fig. 2b is less than the expected value of 67 nm. This indicates that this lamella is not parallel to the foil plane but at an angle. Assuming a d period of 67 nm, one obtains an angle between the fibril and the foil of 42° . Assuming that the fibrils on the left are perpendicular to the foil axis, the angle between the two lamellae is $(90-42) = 48^\circ$, confirming that the angles between adjacent lamellae vary.

AFM corroborates the TEM characterization and provides additional detail (Fig. 2c and d). Since the fibrils were peeled from the scale and the AFM tip provides an accurate representation of the surface, the d banding observed is the actual value of 67 nm. This is shown clearly in Fig. 2d. The three-dimensional rendition in the lower left hand corner of Fig. 2d shows that the fibrils are indeed cylindrical.

3.2. Protective function of the entire scale against penetration

The penetration of the scale by a piranha tooth (Fig. 3a and b) is severely hampered by the mineral layer, amounting to a ~ 1 mm deep indentation into the scale under a 1 kgf load applied for 10 min, with relatively minimal damage around the indent (shown in Fig. 3a). The surroundings of the indent were damaged more seriously due to the severity of load concentration and serrations on the edge of the teeth (shown by the detail in Fig. 3a). Two orientations of collagen fibrils were exposed at the wall of the indent (circular line in Fig. 3a) as the external layer became damaged (Fig. 3b); the fibrils are delaminated, fractured and change orientation (Fig. 3b); these processes permit the internal collagen layer to dissipate energy during penetration.

Fig. 3c and d shows the damage from penetration of a microindenter in the collagen layer (without the mineral layer). In the vacuum of the SEM, the collagen layer dries and shrinks, and separation between collagen fibers becomes evident, which is not characteristic of the intact, water-saturated scale. Nevertheless, the damage mechanisms around the indentation can still be clearly seen. Different orientations of collagen fibers are observed at different levels. The potential indentation region is marked by the dashed square in Fig. 3c. Collagen fibers are bent and fractured through the action of the indenter, but the overall effect of the different lamellar layers is to localize damage to the immediate vicinity of the indenter, without propagation throughout the scale. The failure of the collagen fibers is preceded by deflection, stretching and necking, characteristic of significant inelastic deformation prior to failure (Fig. 3d). As a result, crack initiation and propagation are avoided.

3.3. Tensile mechanical behavior of scales

Fig. 4 shows the uniaxial tensile response of the entire scales and the collagenous lamellar layer (after removal of the external mineralized layer) in the longitudinal and transverse orientations defined in Fig. 1b. Three specimens were tested in each condition. Additionally, the tensile response of the collagenous lamellar layer, including the load–time curve and the behavior of collagen fibrils

Table 1

Uniaxial tensile testing results for the entire scale and for the collagen layer, loaded in the longitudinal and transverse orientations.

	Longitudinal orientation		Transverse orientation	
	Entire scale	Collagen layer	Entire scale	Collagen layer
Young's modulus (GPa)	0.86 \pm 0.32	0.47 \pm 0.25	0.21 \pm 0.02	0.21 \pm 0.03
Ultimate (engineering) tensile stress (MPa)	23.6 \pm 7.2	36.9 \pm 7.4	14.2 \pm 1.1	21.8 \pm 2.4
Engineering strain at maximum stress	0.08 \pm 0.07	0.14 \pm 0.06	0.12 \pm 0.01	0.18 \pm 0.01
Energy dissipation (MPa)	1.47 \pm 1.08	3.12 \pm 0.98	1.07 \pm 0.08	2.53 \pm 0.40

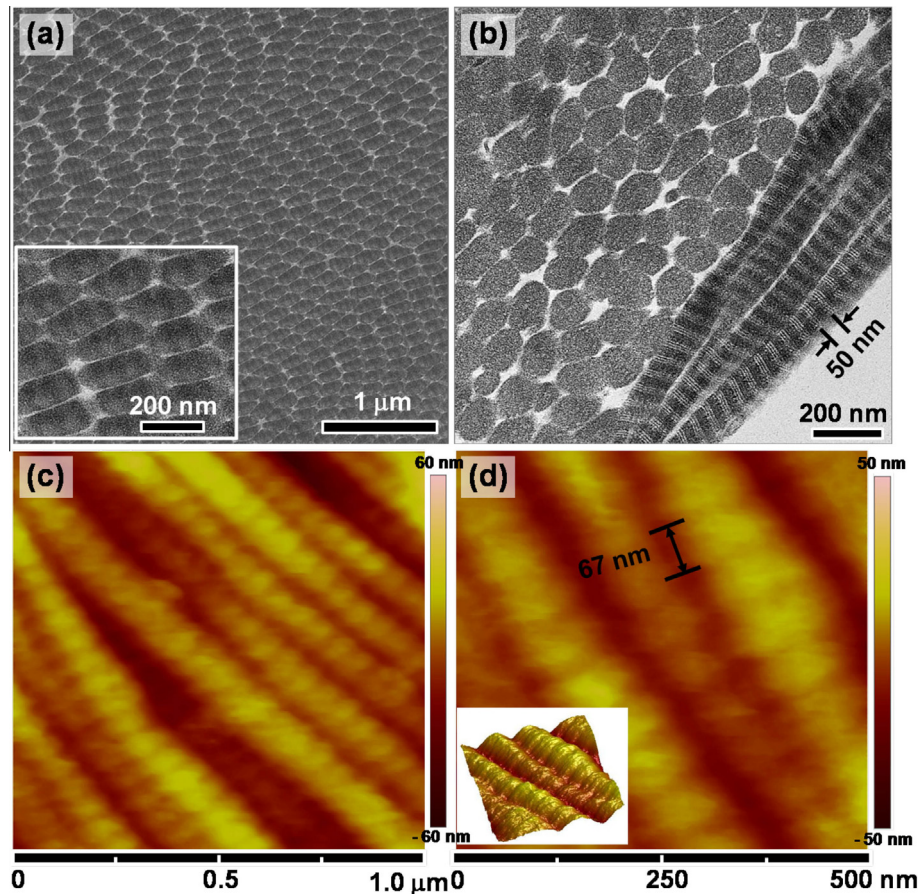


Fig. 2. Characterization of collagen by (a, b) TEM and (c, d) AFM. (a) TEM image of the foil plane inclined to fibril axis; the smallest dimension of the fibrils is ~ 100 nm. (b) Two orientations of fibrils are shown: on the right-hand side the fibrils are inclined to the foil plane showing d bands of 50 nm (projected value of 67 nm), on the left-hand side the fibrils are close to being perpendicular to the foil. Note the regular fibril diameter, ~ 100 nm. (c) AFM of parallel fibrils. (d) Higher magnification AFM showing 67 nm d bands and tridimensional rendition of fibrils in the insert.

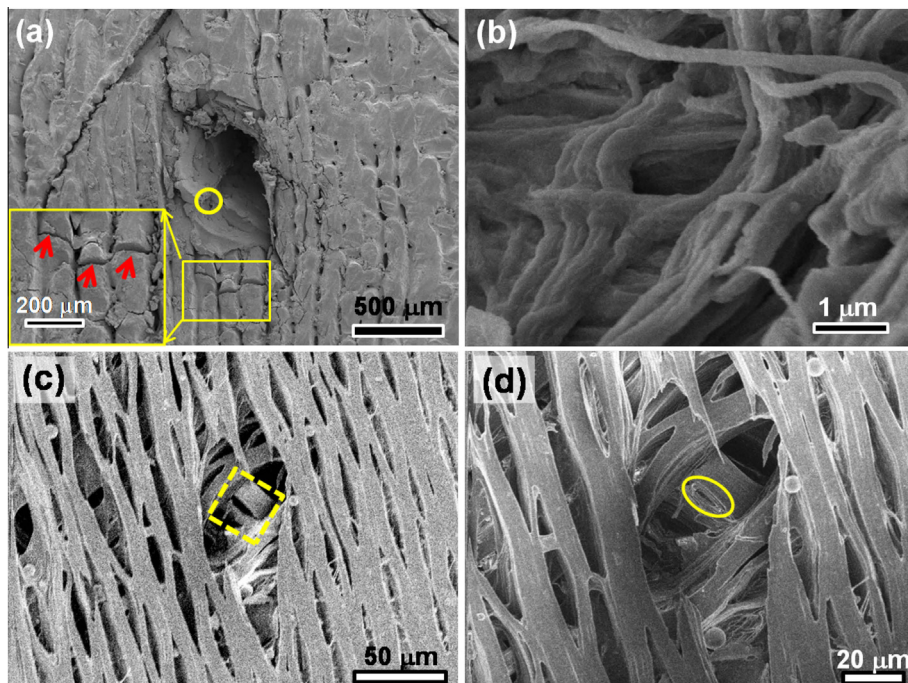


Fig. 3. (a, b) Penetration of the scale by a piranha tooth from the outer layer towards the inner layer. (a) Entire penetration indent of the scale; cracks were deflected and arrested at orthogonal cracks caused by fracture of the mineral (red arrows), with more serious damage at the corner of the penetration. (b) Collagen fibrils exposed from the damage of the mineral observed from the circled region in (a). (c, d) Penetration of collagen layers by a microhardness indenter; the original indentation is shown by the dashed square; the fibers are displaced but the gaps on the samples are due to dehydration in the vacuum of the SEM, the ellipse in (d) is the region that was produced by the indenter tip.

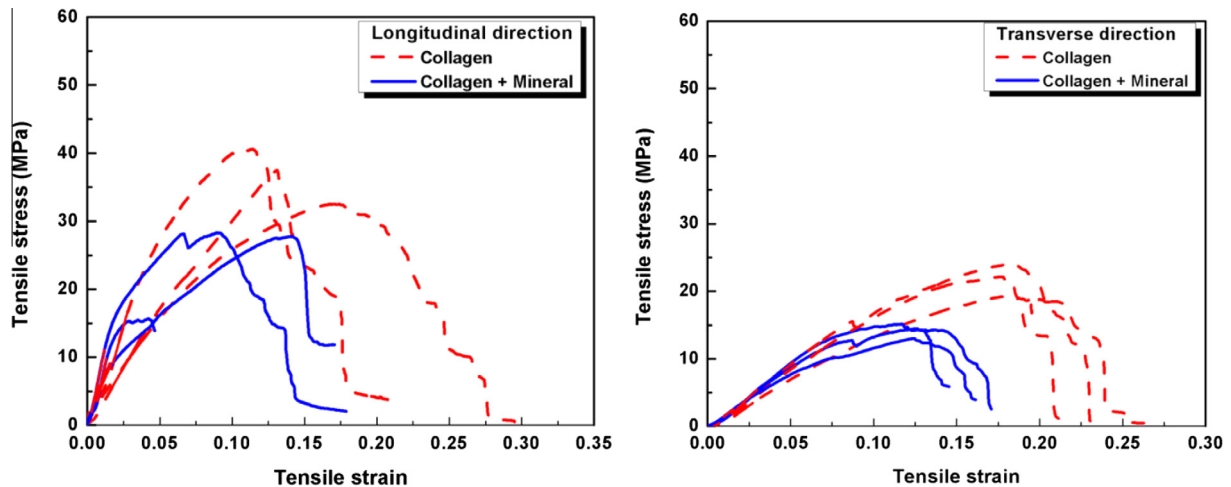


Fig. 4. Uniaxial tensile stress–strain curves of the scales (solid lines) and of the collagen layers with mineral removed (dashed lines). Tests were performed at a strain rate of 10^{-2} s^{-1} in the (a) longitudinal and (b) transverse directions, as defined in Fig. 1.

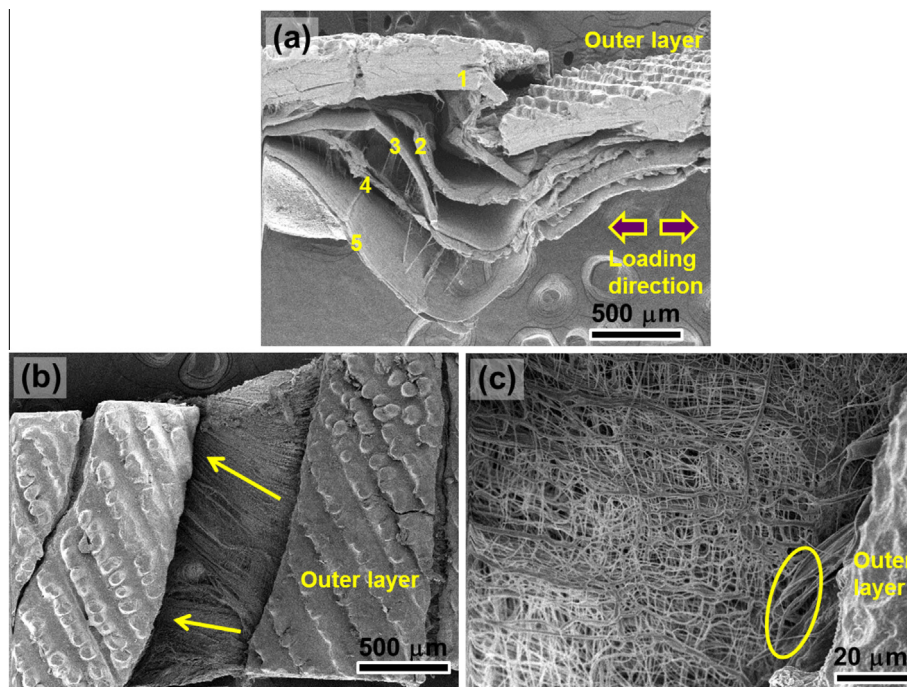


Fig. 5. SEM images of tested tensile samples: (a) side of a tensile sample of the scale after mineral failure, five lamellae were delaminated from each other labeled from 1 to 5; (b) the top view of the tensile sample shows the mineral domains separated and the collagen exposed (the arrows show two orientations); (c) fractured collagen fibrils, some of which are close to the mineral layer (shown by the ellipse); the collagen fibrils beneath are delaminated and separated.

in different lamellae, was also determined in the SAXS tests, as discussed in Section 3.5. Based on these tests, the following results can be deduced (deformation mechanisms are discussed in Section 3.5):

- a. The strength of the scales is $\sim 50\%$ higher in the longitudinal than in the transverse orientation, indicating that the lamellae are not distributed to produce in-plane isotropy. Contrary to our results, Zhu et al.'s [24] measurements on striped bass show that specimens with an orientation of 45° or 90° to the longitudinal axis of the fish have higher strengths ($\sim 60 \text{ MPa}$) than the 0° orientation ($\sim 40 \text{ MPa}$). This shows that in-plane anisotropy exists in both the cycloid and ctenoid scales. The strength of the scales including the

mineralized layer in the longitudinal direction (23.6 MPa) is considerably higher than in the transverse direction (14.2 MPa) shown by Table 1.

- b. Removal of the external and highly mineralized outer layer increases the tensile strength of the scale. Although this appears to be counterintuitive, it results from several factors. Most importantly, the external layer is designed for compressive strength; being highly mineralized it is brittle and weak in tension. This is consistent with the SEM images in Fig. 5a and b, which show the tension samples in the transverse orientation after failure; the external layer fractures in a brittle manner either through the attack of a piranha or other circumstances, leading to tensile stresses in that layer (Fig. 5a: mineralized layer is labeled 1 and has a thickness of

~280 μm). Such a fractured layer cannot carry any load, which leads to a decreased strength of the entire scale resulting from the increased cross-section compared to the collagen-only samples. Additionally, due to failure of the outer mineral layer, there is very likely significant damage evolution in the form of delamination and separation of fibers in adjacent lamellae prior to failure of the entire scale (Fig. 5c). Fig. 5a shows five lamellae, marked by the numbers 1–5 (thickness 50–60 μm), throughout the thickness of the scale; the lamellae close to the mineral layer might be hindered in their ability to rotate upon tensile loading, which could reduce the strength of the collagen layer. The function of the mineralized layer can be rationalized through

a comparison with modern armor, which consists of a hard face supported by a tougher foundation. The hard layer operates under compression and therefore resists projectile penetration, whereas the backing, being tougher and more flexible, ensures the integrity of the structure.

- c. The tensile behavior of the collagen layers was examined in the wet condition; the solid plots in Fig. 4 represent the tensile stress–strain curves of these collagen layers. It is interesting to note that the collagen layers in the transverse orientation have the same modulus as the scale, but a larger strain at failure, suggesting that when the scale is dried and bent in this orientation, cracks are not easily generated. Table 1 shows the tensile properties of the scale in the two

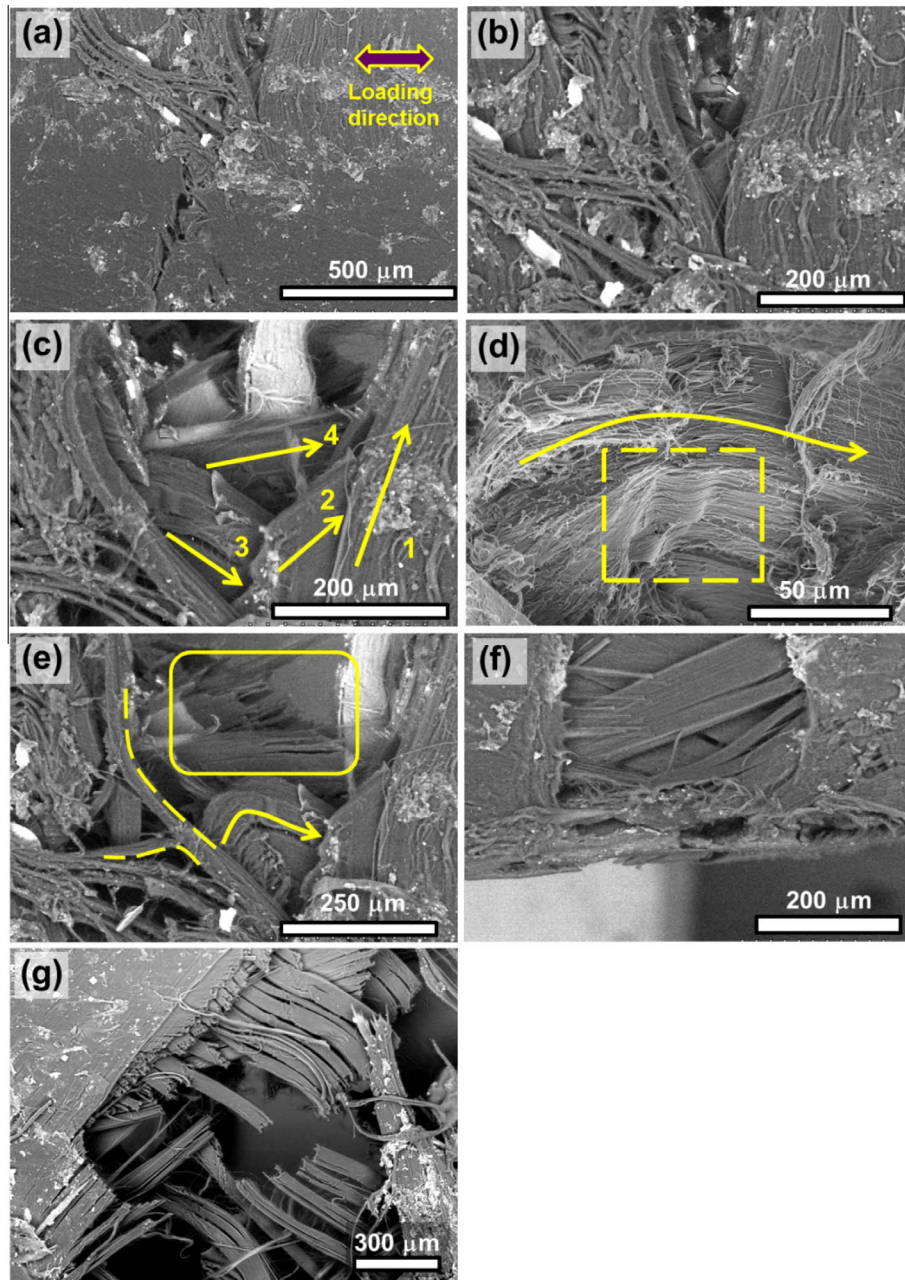


Fig. 6. Damage propagation in the collagen layers containing a notch: (a) initial configuration at the onset of loading (the notch is in the center of the image with the notch root pointing towards the bottom); (b) the collagen fibrils separate when the samples are being loaded; (c) four orientations of lamellae are being exposed (shown by the individual arrows); (d) the collagen fibrils bend and stretch as shown by the arrow, some of the collagen fibrils were relaxed when the test stopped; (e) the lamellae and collagen fibrils bend (arrow), delaminate (dashed lines) and fracture (square); (f) continued loading characterized by sliding of the lamellae indicating the absence of significant crack propagation; (g) damage in unnotched specimen tested under the same configuration up to failure.

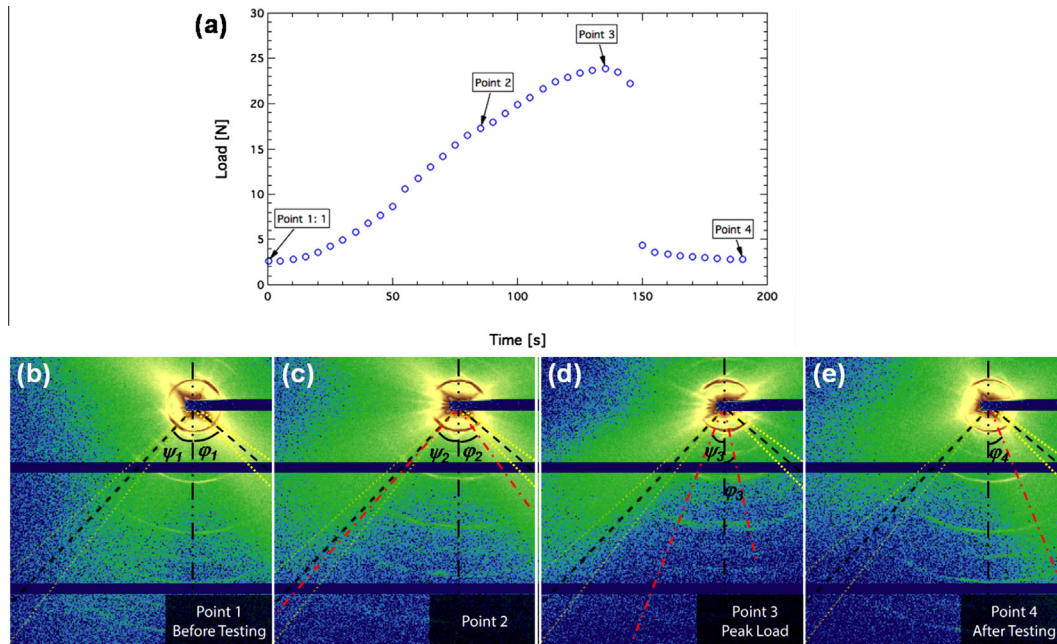


Fig. 7. SAXS results of arapaima collagen layer under tensile load. (a) Tensile load–time plot, SAXS images were taken at four points: (b) before testing, (c) during testing, (d) at the peak load, (e) after testing. Note that the black dashed line shows the average angle (orientation) of the lamellae in scale before tests, the red short dash–dot line shows the average angle (orientation) of the lamellae during the *in situ* tension tests, orientation change of the lamellae can be observed easily from the difference between black dashed line and red short dash–dot lines in the individual (c–e).

orientations (longitudinal and transverse). As noted above, the collagen-only samples have a higher strength and display higher uniform strain and energy dissipation (as measured by the area under the stress–strain curves) than the complete scale samples.

3.4. Tensile behavior of the lamellar structure in notched and unnotched specimens

Fig. 6 shows crack propagation in a specimen in which the highly mineralized external layer was removed, i.e. exclusively in the lamellar structure of the collagen. A crack starting from the root of the notch is forced to grow under the action of tensile forces with a direction marked in Fig. 6a. At the onset of crack propagation, collagen fibers in different orientations are exposed and show different damage mechanisms (Fig. 6b and c). The top lamella (labeled 1), with an orientation of the collagen fibrils perpendicular to the loading direction, undergoes separation by splitting. The fibers in the other underlying orientations (labeled 2, 3 and 4) are more closely aligned to the loading direction and are stretched and rotated. The fibrils (labeled 2) close to the split lamella (labeled 1) fracture; however, some lamellae (labeled 3 and 4) are only stretched and rotated (Fig. 6c and d). Some collagen fibrils are bent (arrow in Fig. 6d) by the loading and then relaxed and curved back after unloading or fracture (dashed square). The full range of failure mechanisms, specifically delamination, tensile failure, rotation and bending of fibrils, is shown in Fig. 6e. Due to the high flexibility of the collagen lamellae and excessive failure distribution upon tensile loading tests, samples showed large crack-tip openings but negligible crack propagation. Failure mechanisms of samples with a highly mineralized external layer occurred in a comparable manner to samples without an outer layer and are hence not shown individually. Fig. 6e and f shows the progression of damage without any appreciable increase in the crack length.

The individual deformation process and mechanism occurring in a tensile test from an unnotched sample are additionally shown

in Figs. 6g and 7 using the SAXS method. The splitting of the individual layers, their rotation and separation as well as curling of fractured fibrils can be also seen. This demonstrates that the processes of fibrillar rotation, stretching, delamination and reorientation occur in the entire specimen being subjected to deformation.

3.5. SAXS, diffraction and mechanisms of collagen reorientation

Synchrotron radiation has been used previously to measure the reorientation and elastic stretching of collagen in bone [35,36] and human aorta [37,38]. The current results for *Arapaima gigas* confirm the previous analyses but importantly introduce another element: the organized reorientation of the lamellae of parallel collagen fibrils, a process that is significantly different from that in other biological systems and which results from the Bouligand-like structure.

SAXS tests were performed on the inner collagen layer to reveal and quantify the mechanisms of collagen lamellae deformation and reorientation during tensile straining, which are essential to the understanding of the resistance of the scale to external loading. The procedures and detailed results are described by Zimmermann et al. [27]; only the principal mechanisms are introduced here to highlight the modes of deformation. The uniaxial load–extension curves on the collagen layer, taken during the SAXS experiment, show an approximately linear response, in agreement with the tensile results of Fig. 4. As the lamellae within the scale's inner collagen layer have a Bouligand-type structure, each lamella will have a distinct orientation that is visible in the two-dimensional X-ray diffraction pattern. Thus, the strain and orientation of the collagen fibers in each lamella can be measured during tensile deformation; the reorientation is clearly visible through the change in the orientation of the arcs with respect to the tensile axis and the strain can be measured by the change in the arc's radial position with respect to the beam center. Four points are marked in the tensile load–time plot (Fig. 7a) measured where SAXS observations were recorded (Fig. 7b and e, respectively), corresponding to points 1–4: before testing (point 1), during testing (point 2), at the peak load (point

3) and after testing (point 4). In the SAXS images, as the collagen fibrils rotate, the angle of the arcs on the SAXS spectra changes; the dash-two dots lines show the tensile loading direction, the dashed lines show the original orientations of collagen fibrils before testing and the dash-dot lines show the changed orientations of collagen fibrils after the testing starts.

Before testing, two main orientations of collagen make angles of $\psi_1 = 43^\circ$ and $\varphi_1 = 49^\circ$ to the tensile loading direction (Fig. 7b). During tensile testing, the collagen fibrils rotate towards the tensile direction (Fig. 7c) such that $\psi_2 = 41^\circ$ and $\varphi_2 = 35^\circ$. At the peak load, the two fibril orientations rotate closer to the tensile direction: $\psi_3 = 21^\circ$ and $\varphi_3 = 12^\circ$. At this point, the SAXS patterns (arcs) of collagen fibrils in two orientations almost meet each other (Fig. 7d). After sample failure (Fig. 7e), one orientation of collagen fibrils releases back to a larger angle ($\varphi_4 = 22^\circ$), whereas the other fibril orientation can no longer be seen, which implies possible fracture.

Such rotations of collagen fibrils both towards and away from the loading direction observed in the *in situ* SAXS experiments can be attributed to two principal mechanisms, as follows.

- Collagen fibril rotation towards tensile loading through interfibrillar shear and elastic stretching (Fig. 8a): when a bundle of collagen fibrils aligned in the same orientation rotate towards the tensile direction, the interfibrillar hydrogen bonds break so that the collagen fibrils can experience shear strain. Fig. 8a shows the collagen fibrils rotating toward the tensile direction ($\psi_1 \leq \psi_0$), as well as their elastic stretching, leading to a tensile strain $\varepsilon = (L_1 - L_0)/L_0$, where L_0 is the initial projected fibril length, L_1 is its stretched length projected in the tensile direction and d_0 and d_1 are the spacing characteristics of collagen fibrils (d period) before and after stretching, respectively.
- Orientation of collagen fibrils change due to formation of an interfibrillar gap (Fig. 8b): when a lamella with collagen fibrils makes a large angle with the loading direction, the fibrils can separate, allowing a gap to open up; thus, regions with one at a larger angle ($\psi_1 > \psi_0$) and another at a smaller angle to the loading direction ($\psi_2 < \psi_0$). In addition to the

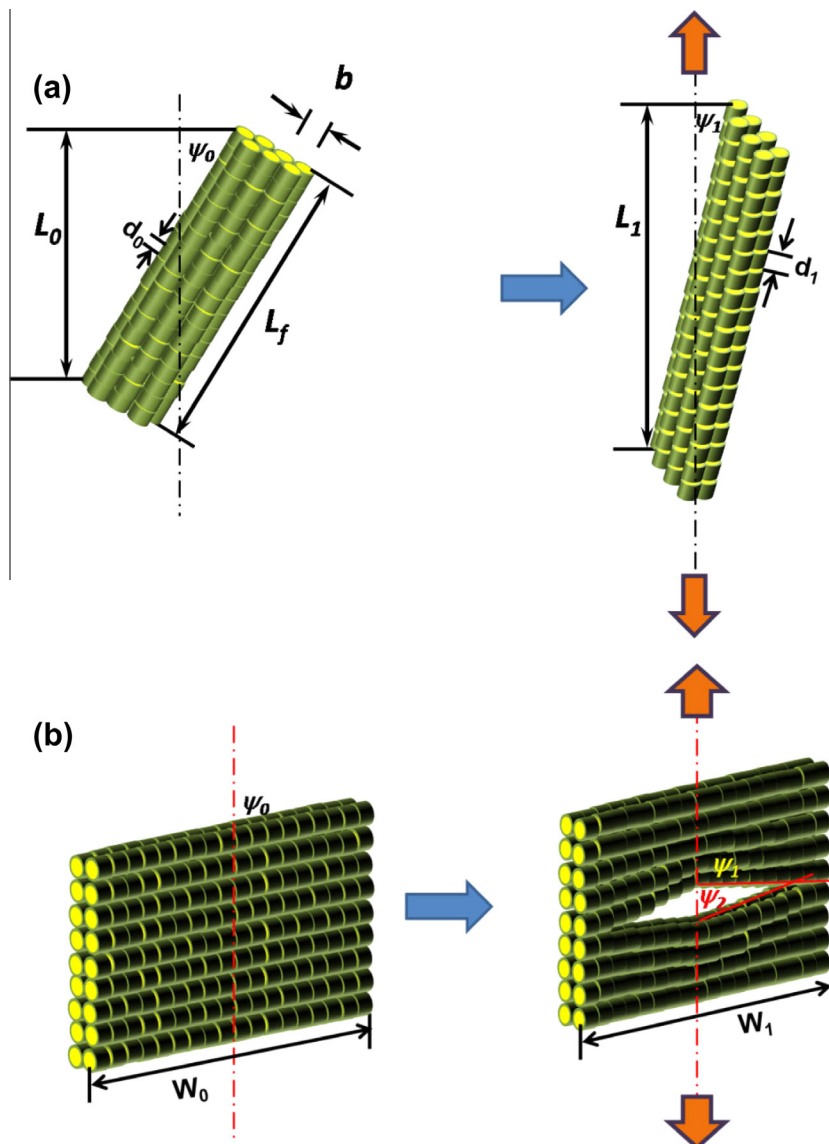


Fig. 8. Mechanisms of collagen fibril deformation under tensile load. (a) Rotation through interfibrillar shear and elastic stretching of collagen fibrils. Fibril stretching and interfibrillar sliding decrease the angle between the fibrils and the loading direction from ψ_0 to ψ_1 . The initial d period along the fibrils is equal to d_0 ; after extension, this d period increases to d_1 as the overall length increases from L_0 to L_1 . (b) The tensile opening of interfibrillar gaps acts to change the angle, at the extremities of the fibrils, from ψ_0 to ψ_1 and ψ_2 ; note the change in width from W_0 to W_1 that can cause Poisson contraction (adapted from Ref. [27]).

above mechanisms, Zimmermann et al. [27] consider “sympathetic” fibril rotation, which can rotate the fibrils toward the tensile axis.

In summary, for collagen fibrils aligning at a small angle to the tensile direction, fibril rotation can easily occur towards the loading direction, whereas for fibrils at a larger angle this rotation is far more difficult. Both rotation toward and away from the tensile axis can be induced.

The rotation of the lamellae toward the tensile axis can be calculated as a function of the total applied tensile strain when the initial angle ψ_0 is small. Two components of strain are considered. The total strain is obtained from the elastic stretching of fibrils (ε_{el}) associated with the increase in d -period of collagen and from the fibril rotation (ε_r) that is the result of interfibrillar shear:

$$\varepsilon_t = \varepsilon_r + \varepsilon_{el}. \tag{4}$$

The strain due to the rotation of the fibrils (involving interfibrillar shear and ignoring the fibrillar stretching) is directly obtained from Fig. 8a; it is equal to:

$$\varepsilon_r = \frac{L_1 - L_0}{L_0} = \frac{\cos\psi_1}{\cos\psi_0} - 1 \tag{5}$$

The elastic strain, shown by the increase in the d period from d_0 to d_1 ($d_1 > d_0$), also has to be expressed in the tensile direction, considering the fibril length L_f and its initial and deformed values, L_{f0} and L_{f1} :

$$\varepsilon_{el} = \frac{L_{f1} \cos\psi_1 - L_{f0} \cos\psi_0}{L_{f0} \cos\psi_0} = \frac{L_{f1} \cos\psi_1}{L_{f0} \cos\psi_0} - 1 \tag{6}$$

The elastic stretching of the collagen fibers follows the linear equation, where E_f is the elastic modulus of the fibrils, estimated to be 2 GPa by Gautieri et al. [34]. Thus:

$$\frac{L_{f1} - L_{f0}}{L_{f0}} = \frac{\sigma}{E_f}. \tag{7}$$

Substituting Eq. (7) into Eq. (6) and this result with Eq. (5) into Eq. (10) gives the strain in the collagen fibrils:

$$\varepsilon_t = \frac{\cos\psi_1}{\cos\psi_0} \left(2 + \frac{\sigma}{E_f} \right) - 2. \tag{8}$$

The constitutive behavior for an assembly of collagen lamellae has been expressed, as a function of strain rate, though a viscoelastic relationship [22]. The one commonly used for biological materials is called the Ramberg–Osgood equation [39]; it was

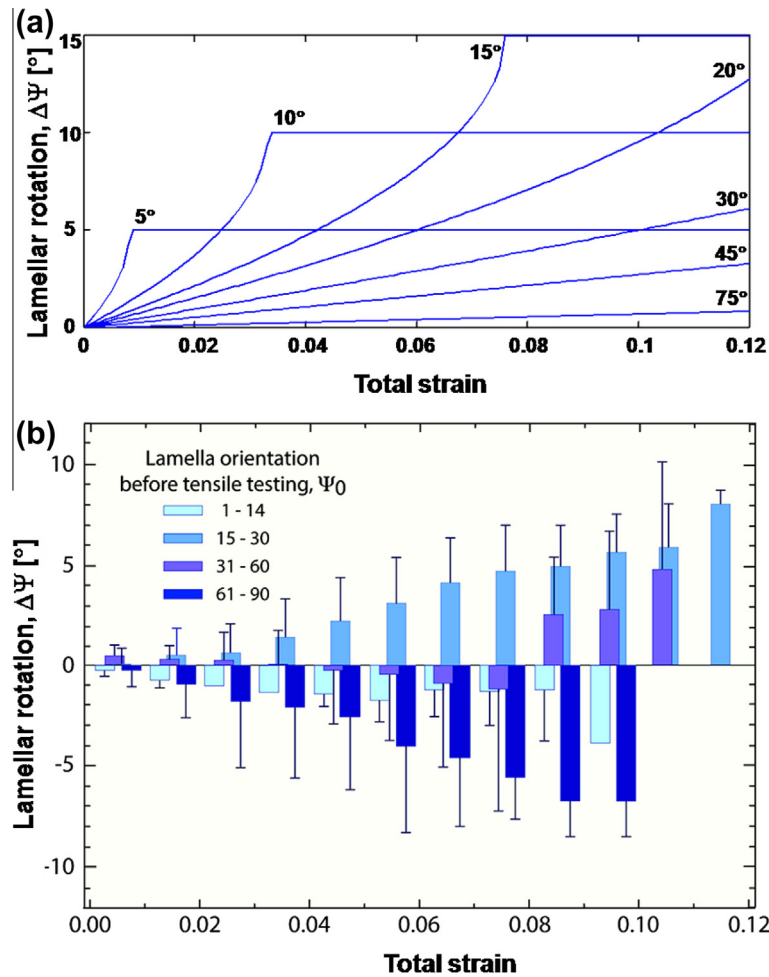


Fig. 9. Nature of the rotation and deformation of the lamellae during tensile loading of the scales. Positive values of the angle $\Delta\psi$ represent rotation of fibrils toward tensile axis and negative values rotation away from tensile axis. (a) Changes in the angles of collagen fibrils with the tensile axis are shown from analytical calculations. Note that when $\Delta\psi$ reaches the value of the initial angle ψ_0 , the rotation stops because the alignment with the tensile axis is reached (at strains ~ 0.01 for 5° and 0.035 for 10°). (b) Changes in the angles of collagen fibrils with respect to the tensile axis measured from small-angle X-ray diffraction patterns; the latter were taken from Ref. [27].

originally developed for metals but its use in the viscoelastic response of bone is well recognized [40]. It describes the strain-rate sensitivity m of the elastic modulus E (which is expressed as σ/ε):

$$\frac{\sigma}{\varepsilon_f} = C \left(\frac{\dot{\varepsilon}}{\dot{\varepsilon}_0} \right)^m, \quad (9)$$

where C is an experimentally measured parameter, $\dot{\varepsilon}$ is the strain rate and $\dot{\varepsilon}_0$ is a normalization parameter, which we can assume to be equal to 1 s^{-1} . Indeed, Lin et al. [22] demonstrated that the arapaima's collagen exhibits a high value of $m \sim 0.26$, with the constant $C = 1.5 \text{ GPa}$. Substituting Eq. (9) into Eq. (8), we obtain:

$$\varepsilon_t = \frac{\cos\psi_1}{\cos\psi_0} \left[2 + C \left(\frac{\dot{\varepsilon}}{\dot{\varepsilon}_0} \right)^m \frac{\varepsilon_t}{E_f} \right] - 2, \quad (10)$$

which is expressed in a format of the changing orientation of the collagen fibrils similar to that used by Zimmermann et al. [27], gives:

$$\psi_1 - \psi_0 = \arccos \left[\frac{\cos\psi_0(\varepsilon + 2)}{2 + C \left(\frac{\dot{\varepsilon}}{\dot{\varepsilon}_0} \right)^m \frac{\varepsilon_t}{E_f}} \right] - \psi_0. \quad (11)$$

Fig. 9a shows the change in collagen orientation given by Eq. (11) vs. the total strain in the range $\varepsilon \sim 0-0.12$. The strain rate used in Eq. (11) was 10^{-4} s^{-1} to match the experimental SAXS results. These values are plotted for different initial values of ψ_0 : 5, 10, 20, 45 and 70°. It can be seen that the fibrils rotate toward the tensile axis for all cases and that rotation stops once the rotation angle reaches the original misorientation angle. When all the collagen fibrils align towards the loading direction ($\psi_0 = 0^\circ$), there is no strain contributed by the rotation of collagen fibrils. For small angles ψ_0 , the rotation is more pronounced. The fibrils originally at an angle between 31° and 90° have no chance to rotate towards the tensile axis and either do not change orientation or rotate away.

The results of the SAXS experiments are given in angular ranges: 0–14°, 15–30°, 31–60° and 61–90°. During tensile testing, we can observe collagen fibrils rotating both towards (positive $\Delta\psi$) and away from (negative $\Delta\psi$) the tensile direction (Fig. 9b). The positive values and the rates of rotation are well described by the above analysis.

Negative values of $\Delta\psi$ (rotation away from tensile direction) require mechanisms not incorporated into the analysis. Three principal mechanisms can be considered: the splitting of lamellae creates regions of separation, causing changes in the angle ψ of the lamellae to the tensile direction, as shown schematically in Fig. 8b (the changes in the angle can be both positive and

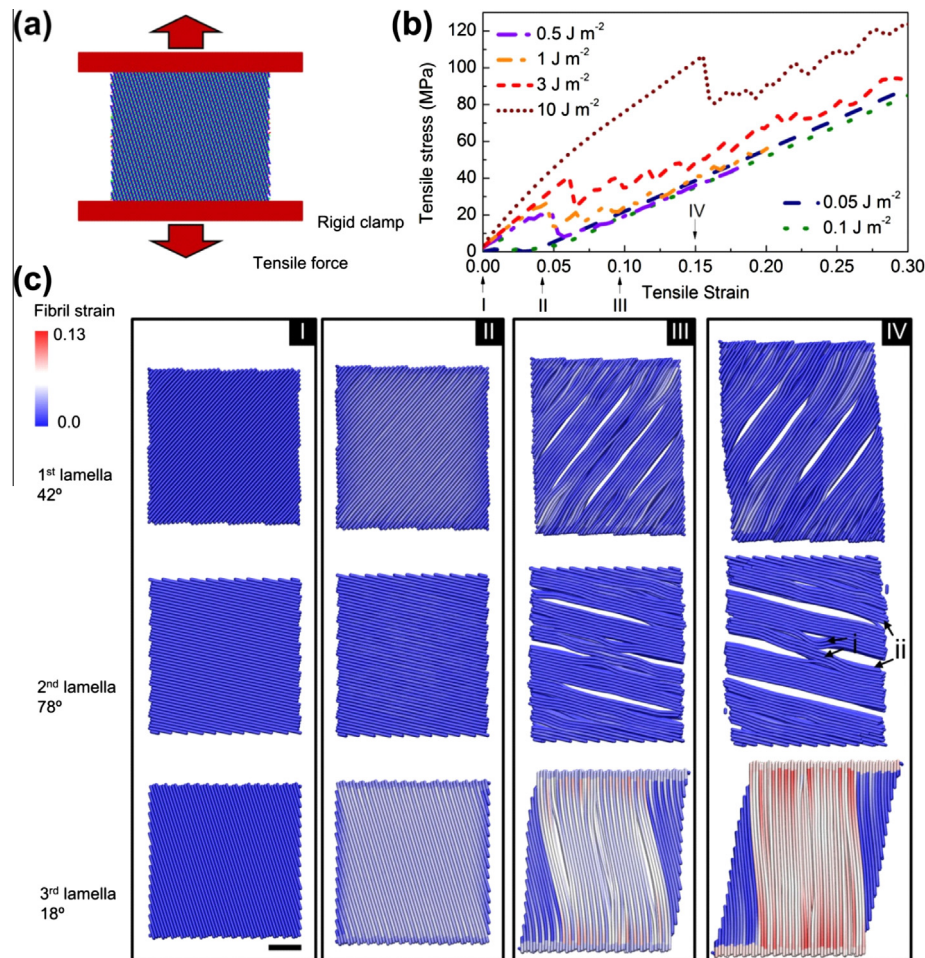


Fig. 10. MD of collagen fibrils in three different directions with initial lamellar angles of 42°, 78° and 18° to the tensile direction. (a) Schematic of the scale model and the loading condition used to investigate the response under uniaxial tension. (b) The stress–strain curves obtained for different surface energies used in model. It is noted that for 1 J m^{-2} the first stress drop is 26.4 MPa, which gives the best approximation to the stress–strain curve of collagen layer and scales as shown in Fig. 4. (c) Simulation snapshots of the deformation of each lamella in the scale model, taken at different strains as noted in panel (b) at increasing strains: (I) strain = 0, (II) strain = 0.04, (III) strain = 0.09 and (IV) strain = 0.15. Collagen delamination (noted by ii) and bridging (noted by i) are marked for lamella 2 at strain = 0.15 (IV), as illustrated (scale bar: 10 μm).

negative); sympathetic rotation through which a layer is sandwiched between two layers which “drag” it and force it to rotate away from the tensile direction; and contraction by Poisson’s ratio effects, leading to rotation. In summary, the analysis predicts the positive $\Delta\psi$ rotations (rotation towards the tensile direction) fairly well. It should be noticed that the analysis uses independent values; no parameters extracted from the SAXS experiments were employed.

3.6. MD calculations on collagen layers in tension

In order to confirm the rotation of the lamellae and individual fibrils observed by SAXS and *in situ* SEM tensile experiments, the process was modeled at the fiber level. MD calculations of the tensile deformation of the collagen layer with three orientations of collagen lamellae (42° , 78° and 18°) were performed to model the mechanisms of deformation. Periodic conditions are applied to the out-of-plane direction and thus each collagen lamella is sandwiched between two other lamellae with a 60° angle with each of them. This configuration is built according to the experimental observations as the angle between two neighboring lamellae is measured to vary between 35 and 85° [22]. An earlier study had reported $\sim 60^\circ$ but this was an oversimplification [23]. We use this model and apply uniaxial tensile force by fixing a single layer of coarse-grained beads at each edge of the pulling direction and controlling their displacement quasi-statically, as detailed in Section 2.8 (Fig. 10a). The Young’s modulus used to model the collagen fiber is 2 GPa, which is an average estimate from collagen microfibrils from both computational and experimental approaches [34]. It is noted that the inter-fiber interaction energy depends on surface chemistry (in terms of protein sequence and mineral composition), degree of hydration and roughness, and it is difficult to directly measure this experimentally. However, we are able to model this interaction by referring to the results of full atomistic simulations. We model the inter-fiber interaction by using the surface energy of tropocollagen molecules, which is measured to have a range from 0.05 to 3 J m^{-2} in simulations with different constraints [41].

Using the present model, we obtain multiple stress–strain curves by using different surface energies, as shown in Fig. 10b. These calculated curves have a typical saw tooth shape, with the stress first increasing linearly with the increasing strain at small deformation followed by significant stress drops before increasing again. The peak stress before the drop region depends on the surface energy; as the surface energy increases, the stress increases, because the bonding between adjacent collagen microfibrils increases. In Fig. 10b, it can be seen that the peak stress increases to over 100 MPa when surface energies are increased to 10 J m^{-2} . For 1 J m^{-2} , the peak stress is equal to 26.4 MPa, which is close to the magnitude of the experimental peak stress of fish scales, as shown in Fig. 4. We therefore use this surface energy to model the inter-fiber interaction. By comparing the simulation snapshots shown in Fig. 10c, we find that during the linear and yielding region, all the collagen fibers in the model retain the orientation of the initial configuration and the deformation strain is homogeneously distributed in the fibers. The first stress drop corresponds to the appearance of collagen delamination within the material; this is an important mechanism for releasing the deformation energy stored in fibers, since it creates microcracks along the fibers. For smaller surface energies (e.g. 0.1 and 0.05 J m^{-2}), the simulations show neither a significant stress drop nor collagen delamination. The last region of the bumpy but upward trend is principally caused by the alignment of the fibers in the third lamella; stretching of those aligned fibers requires larger stress. It is noted that the stress can increase significantly in this region because we assume that the collagen fibers are purely elastic and

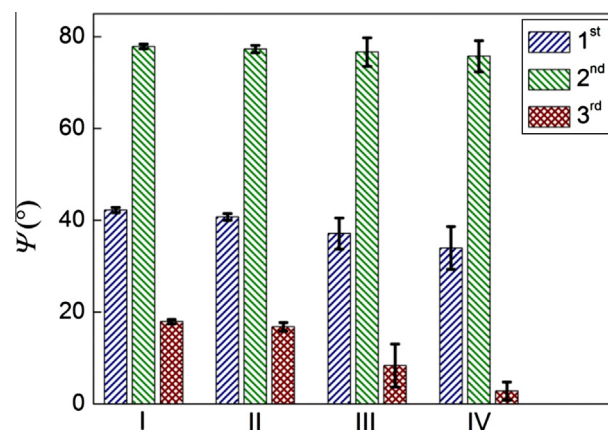


Fig. 11. Simulation results showing evolution of angles ψ of the collagen fibers in each lamella for increasing strains ($I < II < III < IV$). The three layers have configurations shown in Fig. 10c. The four strains used here to scan the angle are 0, 0.04, 0.09 and 0.15 for strains I–IV (same as that used in Fig. 10c), respectively. The variation in the angles increases with strain (marked by bars on top of each bar plot). Lamella 3 shows a significant decrease in angle with increasing strain due to rotation of fibers toward the tensile axis.

non-breakable, which simplifies the mechanics but quickly enables us to see the strain distribution and mechanisms of the collagen delamination and bridging. An improved model needs to incorporate the entire force–extension curve of the collagen fiber; this can be obtained in future from either micro tensile test experiments or large-scale modeling of collagen fibers using molecular simulations [42]. Fig. 10c shows the change in orientation and delamination of the lamellae as a function of applied tensile strain. Four values were used for comparison: 0, 0.04, 0.09 and 0.15. They are labeled I, II, III and IV, respectively. The delamination is more prominent for the first and second lamellae, which have initial angles of 42° and 78° . In the regions at the ends of the delaminations, the angles change significantly. This is indicated by arrows for the second lamella at strain IV. This rotation at the end of the delamination supports the mechanism derived from experiments, as shown in Fig. 8. For the third lamella, which makes an initial angle of 18° to the tensile axis, the rotation towards the tensile direction reduces the angle.

The average value and standard deviation of the angles between the fibers and the tensile direction in each collagen lamella was computed; results are summarized in Fig. 11. It is shown that for the lamella with the largest initial angle (78° , second lamella), the average value stays at a constant level, which agrees with the analytical calculations (Fig. 9a). The standard deviation of the angle in this lamella increases significantly for large deformation, which can be understood by collagen bridging, explaining how a part of the collagen fibers rotates away from the tensile direction (Fig. 10c). The calculations also show that the lamella with a smaller angle with the tensile direction (third lamella, making an initial angle of 18°) rotates faster than the other lamella (first lamella, with angle of 42°) toward the tensile direction, which also agrees with the analytical calculations (Fig. 9a).

4. Discussion

The arapaima scale has two important functions: (i) it resists penetration and (ii) it redistributes the force from the location of the bite over a much larger area. According to Zhu et al. [16], the whole penetration process can be divided into three stages with progressively increasing penetration depth: stage I is essentially elastic loading, which terminates with cracking of the external mineral layer; stage II involves the penetration of collagen; while in

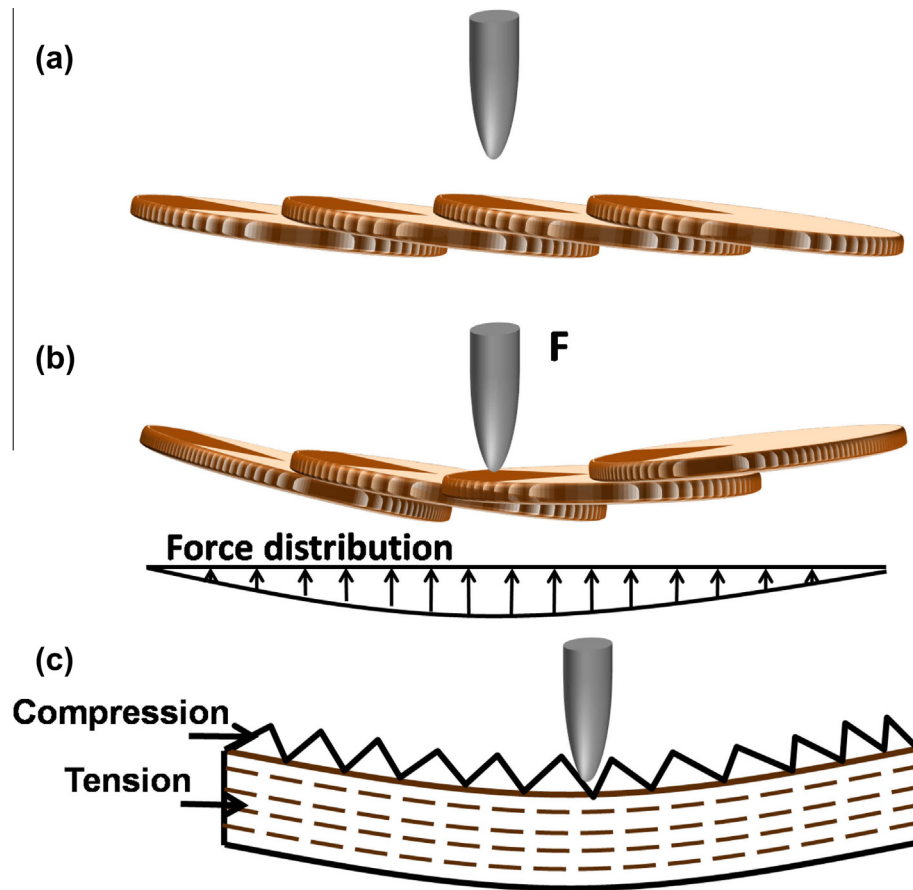


Fig. 12. Protection mechanisms of the arapaima's scales against the penetration of a tooth: (a) schematic configuration of the scale overlap (imbrication); (b) flexing and redistribution of compressive stresses on the tissue; (c) flexure of scale leading to compressive stresses in the mineralized surface layer and tensile stresses in the flexible collagen lamellae.

stage III the tip of the indenter completely traverses the scale. This is indeed what happens in the scale of the arapaimas. Song et al. [25] conducted a finite element analysis of the ganoid scales where they varied tooth and scale parameters and established what effects this had on the mechanical response. One important conclusion is that the radius of curvature of the tip of the tooth had a profound effect on the force necessary to penetrate the scale. This was extended to elasmoid scales by Browning et al. [26] and Varshney et al. [43]. Their conclusion was that the redistribution of loads around the penetration point is critically dependent on the following scale parameters: angle of the scales, degree of scale overlap, composition (volume fraction of the scales), aspect ratio of the scales and material properties (tissue modulus and scale modulus).

Fig. 12a shows schematically how the stresses are redistributed over a large area by the collective action of the scales. This enables the flesh to resist the compression by the application of the bite force. The flexibility of individual scales and their overlap are designed to increase the area of redistribution, as was pointed out by Vernerey and Barthelat [44]. Specifically, the scales are designed to function with the external layer under compressive stresses (Fig. 12b). Indeed, Meyers et al. [23] proposed that the design of scales could lead to a bioinspired flexible ceramic composite and suggested that, in the case of the arapaima scales, the surface ridges minimize the stresses in the ceramic and enable flexing without fracture.

The foregoing results of experiments, characterization and analysis provide an improved understanding of the mechanisms of deformation and damage in the scales. The external mineral layer

is designed to operate under compression; the tensile strength of the scales is actually increased by its removal. This is paradoxical but can be explained by the fact that the mineral does not contribute significantly to the tensile strength. Thus, its effect is strictly to increase the cross-sectional area, thereby decreasing the stresses. A second effect is that the lamellae adjacent to the mineral layer do not benefit from the rotational mechanisms discovered by Zimmermann et al. [27]. Indeed, the mineral layer is designed to operate in compression, as shown in Fig. 12b. The application of a force by a tooth bends the scale in such a manner that the external mineralized layer is put under compression whereas the internal layers are subjected to tension (Fig. 12c). In fact, penetration by a tooth is hampered significantly by this layer, leading often to the fracture of the tooth.

The lamellar structure of the collagenous foundation comprises many layers in a Bouligand-like arrangement. As a tooth penetrates, the layers separation and scale fracture are avoided through interlamellar and interfibrillar sliding mechanisms. The reorientation of the lamellae was observed through *in situ* experiments conducted on pre-notched specimens. These experiments show the sliding and reorientation of the lamellae and the absence of crack propagation. SAXS experiments confirm the change of orientation of the collagen fibrils. These phenomena were modeled by MD; this confirms that at small angles ($\psi < 45^\circ$) the lamellae rotate towards the tensile axis, while for larger values, interlamellar splitting causes local increases in ψ .

The combination of a highly mineralized surface region designed to operate in compression with a flexible foundation,

which can undergo significant tensile stresses, provides mechanical properties that minimize the penetration ability of teeth and can distribute the compressive traction effectively while ensuring significant mobility to the fish. This is accomplished at a modest weight penalty: the scale weight/fish weight ratio is only ~ 0.1 .

5. Conclusions

Arapaima gigas is one of the largest freshwater fish; it is covered with elasmoid scales that act as flexible dermal armor and provide protection, primarily from attacks by piranhas, the principal predators in the Amazon basin. In this study, the penetration response and tensile behavior of the scales were investigated, with particular attention to the mechanisms of damage in the individual collagen layers. This was achieved experimentally through the use of tensile tests on hydrated scales and indentation experiments, analytically by predicting the various mechanisms that retard damage in the scales, and computationally with MD calculations performed to understand the response of collagen fibrils to tensile loading. Based on these studies, the following specific conclusions can be made:

TEM and AFM confirm that the collagen fibrils have diameters of ~ 100 nm and are parallel in one lamella. The d band spacing of the collagen is clearly revealed by AFM and is equal to 67 nm, characteristic of type I collagen.

The scales act to efficiently resist bites from other fish. As a tooth attempts to penetrate the scale, first the brittle external layer is placed under compression and would eventually crack due to the stress concentration and gradients, exposing the collagen layers beneath. As the penetrator enters more deeply into the scale, the collagen fibrils are pushed apart and separated. Since it has to traverse several lamellae with different orientations, no macroscopic cracks are formed. In an extreme case, eventually, individual collagen fibers fail in tension, exhibiting necking which is characteristic of considerable permanent deformation.

We confirm and quantify earlier conclusions by Zimmermann et al. [27], namely that the stretching, rotation, delamination and fracture of collagen fibrils are the principal mechanisms of energy dissipation in the flexible foundation. Under tensile loading, most of the collagen fibrils are stretched and rotate toward the tension direction; the collagen fibrils with a large angle to the loading (closer to 90°) delaminate as portions can rotate away or towards the tensile direction. An analytical model predicts the rotation of the collagen fibers toward the tensile axis; the model is in good agreement with measurements by SAXS.

MD calculations of interfibril separations yield results that match quantitatively the experimental measurements, with interaction energy between microfibrils of 1 J m^{-2} being assumed. The calculations correctly predict the separation of fibrils and angular rotation toward and away from the tensile axis for large angles, and rotation of fibrils toward the tensile axis for small angles. Since only three lamellae are used in the computation, their splitting is accompanied by significant load drops; the drop at a stress of 26.4 MPa corresponds closely to the maximum stress of 30–40 MPa experimentally observed.

Acknowledgments

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Appendix A. Figures with essential colour discrimination

Certain figures in this article, particularly Figs. 1–12 are difficult to interpret in black and white. The full colour images can be found in the on-line version, at <http://dx.doi.org/10.1016/j.actbio.2014.04.009>.

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