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Non-invasive diagnostics in pathological fossils by magnetic resonance imaging

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Abstract

For more than a decade, Magnetic Resonance Imaging (MRI) has been routinely employed in clinical diagnostics because it allows to non-invasively study anatomical structures and physiological processes in vivo and to differentiate between healthy and pathological states, particularly in soft tissue. Here, we demonstrate that MRI can like-5 wise be applied to fossilized biological samples and help in elucidating paleopathological and paleoecological questions: Five anomalous guards of Jurassic and Cretaceous belemnites are presented along with putative paleopathological scenarios directly derived from 3D Magnetic Resonance images with microscopic resolution. These syn vivo deformities of both the mineralized internal rostrum and the surrounding former 10 soft tissue can be traced back in part to traumatic events of predator-prev-interactions. and partly to parasitism. Evidence is presented that the frequently observed anomalous apical collar might be indicative of an inflammatory disease. Finally, the potential of Magnetic Resonance techniques for further paleontological applications is being discussed. 15

1. Introduction

Paleopathological studies can shed light on the ecology of ancient life forms, and they are helpful to evaluate the functional significance of morphological traits (Keupp, 1984 and 1985). Most of the described paleopathological phenomena of fossil hard parts
²⁰ have a synecologial origin, i.e. they arose from interactions between different species (e.g. through predation, parasitism, epizoism). The specific reaction of the organism under attack lead to a characteristic syndrome whose symptoms often include anomalous growth patterns that can still be observed in fossil remains. The resulting degree of alterations in morphology always ranges within specific tolerance limits and allows
²⁵ to estimate the functional relevance of distinct morphological features.

Within a wealth of paleopathological studies, most focus on vertebrates (cf. Tasnadi-

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Kubacska, 1962) but considerable efforts have also been directed towards pathological aspects of invertebrates - namely ectocochleate cephalopods, and ammonites in particular, research on which has a long scientific tradition (e.g. Engel, 1894; Hölder, 1956; Guex, 1967; Bayer, 1970; Keupp, 2000; Hengsbach, 1996; Kröger, 2000; Ke-5 upp, 1984 and 1985). Descriptions and interpretations of anomalies in skeletons of endocochleate cephalopods, in contrast, appear only sporadically (for modern sepiids, see Ruggiereo, 1980; Battiato, 1983; Ward and von Boletzky, 1984; von Boletzky and Overath, 1989; Bello and Paparella, 2002; for fossil squids, see Dietl and Schweigert, 2001). One notable exception are healed fractures in belemnite guards (see references in Keupp, 2002) which, on the one hand, were seen as a proof of the 10 endocochleate nature of the belemnite skeletons and, on the other hand, gave rise to speculations about their function being, amongst other suggestions, a pushing weapon (Naef, 1922) or a digging tool (Abel, 1916). The reason for the different treatment of endo- and ectocochleate specimens has a very practical nature: The shells of ectocochleate cephalopods allow sublethal injuries and other growth disturbances as well 15

- as the subsequent formation of characteristic syndromes of the attacked animal to be diagnosed immediately from anywhere in their posterior parts. Conventional investigations of anomalous belemnite guards, though, require to damage the rostrum in which former growth stages are surrounded by the later calcitic layers. Similar acces-
- sibility constraints apply to other types of specimens whose analysis by such invasive methodology, consequently, leads to considerable damage or even loss of the sample (Siveter et al., 2004). Our study, therefore, was designed to evaluate the potential of a non-invasive technique – Magnetic Resonance Imaging (MRI) – for paleontological applications, particularly in cases of rare anomalous fossils where it is especially desir-
- ²⁵ able to avoid specimen destruction or to make informed decisions about how to sample invasively, where and how much.

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1.1. Magnetic resonance imaging

The concept of Magnetic Resonance techniques and their applications in materials science and biomedical research have elsewhere been described in detail (e.g. Callaghan, 1991; Blümich and Kuhn, 1992; Boesch, 1999; Glover and Mansfield, 2002). In biomineralized samples, they have been employed for bone trabecular structure determinations in osteoporosis patients (Borah et al., 2001), for comparative studies on skeletal evolution (Spoor et al., 2000) as well as for indirect imaging of moldic fossils after artificial (and destructive) filling with water as a contrast agent (Clark et al., 2004), and a short review on their potential for archeological investigations including resins, wood and semi-fossil bones is available (Lambert et al., 2000). Briefly, the spin of the nucleus of each atom possesses an isotope-specific resonance frequency directly proportional to the magnetic field strength it experiences. Thus, if irradiated by radiowaves at or very close to this frequency, a subpopulation of such an isotope can be excited. In a highly homogeneous field, subtle differences between the chemical

- ¹⁵ bonds of different nuclei of the same species will translate into shifts of their resonance frequency, the recording of which is the basis of MR Spectroscopy (MRS). If, moreover, uniform magnetic field gradients are superimposed to the static field after excitation, the signal response which relaxes with two major time constants, T₁ and T₂, will contain frequency-encoded information about the spatial distribution of the excited
- ²⁰ nuclei, which allows for MR Imaging. In our study, only the signal of hydrogen nuclei (protons) within the fossils was detected. Depending on the chemical composition of a sample, on the gradient strengths and on the excitation pulse sequences used for data acquisition, the chemical and spatial information can be separated in MR images. However, for the solid state samples investigated here, the signal intensity obtained for
- each volume element (voxel) mainly reflects its hydrogen content and is insensitive to properties of the chemical bonds that the protons in this voxel are engaged in. Therefore, a clear distinction of signals originating from organic or inorganic compounds of these belemnites is not possible.

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1.2. Architecture and normal growth pattern of belemnite guards

The belemnite guard formed part of the endoskeleton of a squid-like decabrachiate coleoid and consists primarily of low-Mg calcite (Sælen, 1989). Its distal part (rostrum solidum) is massive, the proximal part (rostrum cavum) surrounds the chambered aragonitic shell (phragmocone) that served as a buoyancy apparatus. The rostrum solidum 5 is sometimes distally covered by an epirostrum that often appears hollow due to a presumably high primary content of organic or aragonitic material (Bandel and Spaeth, 1988). The guard is generally thought to have served as a counterweight to the soft body and was situated at the posterior portion of the orthoconic phragmocone, allowing the animal to maintain a horizontal swimming posture (Naef, 1922; Fischer, 1947; 10 Stevens, 1965). The primary anlage of the rostrum was the cone-shaped "primordial guard" within the egg (Hanai, 1953; Jeletzky, 1966; Bandel et al., 1984). During ontogenesis, the rostrum thickened concentrically, and sometimes allometrically, by periodic accretions of radially arranged calcite prisms. The resulting microstructure is characterised by dominating radial calcite fibres with numerous relatively distinct growth

- lamellae that transverse the calcite prisms and become more closely spaced towards the outer margin of the guard. The original texture of the orthorostrum was solid, with an inner porosity of only 10 % maximum (Veizer, 1974). Distinct concentric growth rings that cut the radially arranged crystals reflect phases of episodically reduced mineralisation that temporarily allowed integration of higher percentages of organic matter (Sælen, 1989), most of which has later been replaced by diagenetic calcite, as evidenced by data of stable isotopes (δ^{18} O, δ^{13} C) and cathodoluminescence (Podlaha et al., 1998; O'Neill et al., 2002).
 - 1.3. Paleopathological phenomena
- Deformations of belemnite guards can result either from diagenetic, tectonic and impact events after the death of the organism or, during its life time, from illness or antibiotic interactions (predator-pray relationships, parasitism). Specimens of the latter

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group are collectively referred to as "pathological" ones (Keupp, 2002) and will be in the focus of this study, especially during an earlier, juvenile growth stage.

Pathological guards were described and interpreted for the first time by Duval-Jouve (1841) in sliced rostra with fractures. First attempts to extract evidence for behavioural

- traits of belemnoid animals from pathologic guards were made by Abel (1916) who attributed Duval-Jouve's rostral fractures, rather than to an unsuccessful attack by a predator, to damage experienced when the animal used its posterior end as a digging tool. Consequently, he postulated a benthic way of life for these specimens. However, cuttlebone growth disturbances in modern cuttlefish living in their natural habitat
- are mostly triggered by repair of sublethal injuries as a consequence of unsuccessful attacks of predators including human divers (von Boletzky and Overath, 1989). Correspondingly, it may be expected that many anomalous guards of belemnites reflect the repair of similar damages to the preadult rostrum or its muscular mantle. Nonetheless, it has been shown that besides such exogenous traumatic events remarkable anomalies also resulted from endogenous disturbances of the mantle tissue, including
- presumed cases of parasitism (Radwanska and Radwanski, 2004).

2. Sample description and method

Five belemnite guards from the collection Keupp (PB), housed at the Institut of Geological Sciences, Freie Universität Berlin, Germany, were investigated. Hereafter, they
will be referred to by their respective catalogue numbers. PB246, PB248 and PB249 are *Gonioteuthis quadrata* (Blainville, 1827) guards from the Late Cretaceous (Upper Campanian, ca. 75 Myr) of the marl pit "Alemannia", Höver near Hannover, Germany. PB264 is a cast of a *Neoclavibelus subclavatus* (Voltz, 1830) rostrum from Late Toarcian (Upper Liassic, ca. 182 Myr) of Mistelgau near Bayreuth, Germany. Our study has been performed on the original sample which is with V. Kriegisch, Schonungen, Germany. PB251 is a *Hibolithes jaculoides* (Swinnerton, 1937) rostrum of the Lower Cretaceous (Late Hauterivian, ca. 118 Myr) from the coast of Helgoland Island, Ger-

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many.

The MRI experiments were performed on a BRUKER Avance NMR spectrometer (Bruker, Rheinstetten, Germany) operating at a ¹H resonance frequency of 400 MHz with standard Micro2.5 microimaging equipment and a maximum gradient strength of

⁵ 0.4 T/m. Images were recorded using a standard 3D spin-echo imaging sequence (echo time $T_e = 1.3$ ms, repetition time $T_r = 1$ s if not mentioned otherwise; for details, see Ernst et al., 1997). Each image consists of $128 \times 128 \times 256$ voxels. With typical sample dimensions of the order $10 \times 10 \times 30$ mm³, this normally resulted in spatial resolutions around 100 μ m. The total acquisition time (T_a) ranges between 18 and 93 hours per image, depending on each sample's signal strength. Details are given in the figure legends.

In the image slices, occasional single pixels with high signal intensity but no morphological context are an artifact originating from the Fourier transformation of frequency into spatial information. The apparent loss of signal intensity towards the edges of the Field of View (FOV) is due to the coil geometry employed to send and receive the

the Field of View (FOV) is due to the coll geometry employed to send and receive the radiofrequency signals. The images were visualized and processed with the help of ImageJ (available at http://rsb.info.nih.gov/ij; developed by Wayne Rasband, National Institutes of Health, Bethesda, MD).

3. Results and discussion

All five belemnite guards described here exhibit anomalous growth patterns, the beginning of which – as well as their presumptive endogenous or exogenous origins – are not observable from outside, since subsequent calcitic lamellae cover the original disruption and obstruct the cause for its existence. In contrast, the presumed origins of the pathologies and their subsequent development of growth anomalies could noninvasively be extracted from the specimens by means of Magnetic Resonance (MR) image series. The following anomalies have been recognised and their causality interpreted: 2, 239-260, 2005

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3.1. Sublethal injury with fracture of the rostrum (Cat. PB246)

The complete, 33 mm long and up to 9 mm broad guard of *Gonioteuthis quadrata* shows a zigzag-like deformation (Fig. 1A). The MR image (Fig. 1B) revealed multiple internal fractures of the juvenile guard (Ø 5.1 mm in maximum) distally of the intact alveole. We conclude that, as the result of a probably sublethal bite of an unknown predator, the second fragment has ventrally been dislocated almost perpendicularly to the two neighbouring fragments, while the small final fragment of the distal tip of the rostrum shows only a tiny kink. A presumed fourth fracture cutting the small fragment near the distal tip produced no dislocation. The fractures healed after the dislocated fragments (Abel, 1916) and was then consecutively covered with several calcareous layers. The interim surface of the guard's fragments at the time of the traumatic attack is marked by sharp lines of high signal intensity, interpreted as to represent a temporary interruption of the mineralization which triggered a higher incorporation of organic material.

- ¹⁵ During regeneration, the secretion of new calcareous layers was partly asymmetrical. Particularly inside the knee-like arrangement of the rostral fragments (i.e. dorsally), the thickness of single growth lines is enlarged, and accreted posttraumatic calcitic layers add up to 6.2 mm in thickness, whereas they are reduced on the convex part – only a 0.8 mm thick posttraumatic calcitic coating was added at the ventral part of the knee. A very similar anomalous guard of *Hibolithes subfusiformis*, described by Douval-Jouve
- (1841, cf. pl. 10, Fig. 17 in there) had recovered from a double-fracture by forming such a double-kinked rostrum.
 - 3.2. Sublethal injury which only affected the muscle mantle around the guard (Cat. PB248)
- ²⁵ Two anomalous rostra of *Gonioteuthis quadrata* exhibit an apparent doubling of their pointed apexes. The 60 mm long and up to 9 mm wide guard PB248 developed, along the final 25 mm of the main guard, a second rostral element laterally, beginning with

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a bubble-like bulge of 9 mm length (Fig. 2A). The phenomenon corresponds with an apparent bifurcation of the rostrum in which the two branches – that originally developed separately and ran subparallel to each other – have in fact been connected in a later phase by common growth layers. The MR image (Fig. 2B–E) shows very clearly that the surface of the juvenile guard inside (Ø 7.6 mm) was probably undamaged at the beginning of the anomaly. The irregular growth pattern from which the bulge arose

- at the onset of anomalous growth through subsequent thickening by new calcite layers can be interpreted as a disturbance of the calcite secreting mantle, most probably initiated by a traumatic event. The muscular mantle had in part torn off, thereby isolating
- ¹⁰ a small piece of it. Similar to the formation of a free pearl in mussels where an isolated fragment of the secreting epithelium provides the basis for the construction of the pearl sac (Götting, 1974; Keupp, 1987), this small piece was then the starting point for a second mantle fold during regeneration of the mantle tissue in which an additional rostrum could evolve. The moment of the traumatic event is marked by a sharp growth
- line inside the entire guard, documenting again the phase of regeneration in which the mineralization process was temporarily interrupted and a distinct layer rich in organic matter inserted. Similar anomalous rostra with proliferate growth phenomena by formation of separated mineralisation centres have been described of a Jurassic *Hibolithes* (Keupp, 2002), while guards with two or multiple tips have repeatedly been reported
 (Keupp, 2002; Wundt, 1883; Schwegler, 1939; Finzel, 1963; Schmid, 1963; Miertzsch, 1964; Ladwig, 1993).
 - 3.3. Apical collar as a result of inflammation (Cat. PB249)

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Another phenomenon frequently observed in Late Cretaceous belemnite guards of different taxa but not yet described is the development of a small collar-like rim in dorsal position above the rostral tip. The 75 mm long and up to 11 mm broad guard PB249 appears to have a double tip, owing to the close position of the anomalous dorsal collar to the normal tip of the rostrum (Fig. 3A). Yet it differs significantly from the real double tips described above. The MR image (Figs. 3B and 3C) reveals that the growth of the 2, 239–260, 2005

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anomalous collar starts at the former surface of the guard with no visible damage and continues into a small hollow tunnel with an apical opening. The small tunnel resembles drainage channels for organic fluids of inflammatory portions of tissue. Therefore, this characteristic anomaly of belemnite guards is presumed to have been initiated by
 a local infection or other inflammatory disturbances of the muscular mantle, perhaps

also by settlement of a tiny parasite (see below).

3.4. Disturbance of the guard secreting mantle by presumed parasitism (Cat. PB264)

Bubble-like protuberances of guards as depicted in Figs. 4 and 5 were very probably initiated by implantation of parasites. The 32 mm long rostrum of *Neoclavibelus subclavatus* exhibits a drop-sized bubble elevated up to 5 mm above the guard's normal surface and shows a growth direction towards the anterior part of the animal (Fig. 4A). The vertical and horizontal MR image sequences (Figs. 4B–4D) show a prominent growth line surrounding the juvenile guard at about 2.9 mm diameter. During subsequent thickening of the guard, the successive formation of an irregularly bordered cavity

- ¹⁵ can be observed immediately above the tip of the interimistic surface of the guard. At the beginning and end of the increasing bubble-like anomaly, no signs of a preceding damage of the rostrum are recognisable (cf. Fig. 4D). The sharp concentric growth line drawing the undisturbed juvenile guard is therefore presumed to mark the temporary interruption of an otherwise rather continuous secreting activity of the muscular mantle
- in response to the infection by an unknown parasite. The protuberance of the guard probably results from the successively increasing proliferation of the endoparasite. A 3D model of the guard, as obtained from the MR images, is depicted in Fig. 4E.
 - 3.5. Early disturbance of the guard secreting mantle by presumed parasitism (Cat. PB251)
- ²⁵ A similar syndrome, probably also due to parasitism, is shown in Fig. 5. The surface of the 43 mm long and outside the anomaly 7 mm wide rostrum of *Hibolithes*

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jaculoides is slightly corroded. An ovoid bubble elevates about 6.5 mm above the normal surface near the proximal part of the guard (Fig. 5A). The MR images (Figs. 5B and 5C) suggest that the formation of the anomalous bubble began during an early ontogenetic stage and enabled the formation of an internal irregularly bordered cavity
⁵ which enlarged the middle axis of the rostrum up to 7 mm. The anomaly starts close to the empty alveole from which only the distal part is preserved, probably situated immediately after the (invisible) primordial rostrum. The internal cavernous structure seems to be divided into a proximal ovoid chamber of ca. 2 mm by 5 mm, potentially reflecting the original size of the encysted parasite itself, and a larger distal portion
¹⁰ characterised by proliferous growth of irregular tissue without or with only slight calcitic mineralization.

4. Conclusions

Our study has shown that Magnetic Resonance techniques allow to non-invasively visualize structural details in fossils. Although this is perhaps surprising on its own – fossils notoriously lack typical MRI contrast generators, namely water and soft tissue – our feasibility proof of fossil MRI may be less striking than the observation that ordinary liquid state pulse sequences lead to reasonable image contrast in rock-solid mineralized materials. What is more, solid state techniques, at least in the belemnite samples we investigated, do not reach a comparable signal-to-noise ratio (data not shown). So the question is where the signal comes from. Fortunately, the anomalous samples

- the question is where the signal comes from. Fortunately, the anomalous samples already give some hints: First, signal intensity in the pathological regions is usually higher (cfs. Figs. 2B and 2D) and rarely lower (cracks in Fig. 1B) than in the normal ones, thereby indicating that the syn vivo processes laying the ground for the uneven signal distribution were activated rather than inactivated by the pathological incident.
- This corresponds well with previous reports of higher organic contents in pathologically affected areas (Sælen, 1989). Second, the frequent observations of a hollow (as in PB264 and PB251) rostrum cavum have led to the suggestion that its primary con-

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tent might have been organic (Bandel and Spaeth, 1988). Third, in cases where the innermost layer of the rostrum is not hollow, it gives a higher signal than its immediate surrounding (as in Figs. 1 and 2). While this might equally point at taxonomic differences, it is well compatible with the assumption that the image contrast in the belemnite

- MRI data basically stems from differences in primary content in organic matter which was, perhaps, later partly replaced by diagenetic calcite. Fourth, the compact concentric layering of separate organic and anorganic layers in belemnites could provide an explanation for the presumed good conservation of organic matter within these structures. However, with the imaging parameters used, hydrogen nuclei of crystal water in
- inorganic materials can usually not be distinguished from hydrogen nuclei in organic molecules. This distinction would require additional experiments with different MR parameterization or complementary methods, and for the samples discussed here, the indication of catalogue numbers will facilitate such comparative methodological studies.
- The MR image data obtained from belemnite guards allowed to observe their smallscale three-dimensional growth lamellae and to consistently interpret origins and developments of internal structures in rostra experiencing ontogenetic or anomalous anatomical and physiological alterations. Furthermore, the notion that the apical collar might be indicative of an inflammatory disease illustrates that MRI can contribute to
- the generation or validation of hypotheses on belemnite pathologies. Finally, all traditional paleontological investigations still remain possible after MRI scanning, which is not necessarily true in the opposite case. Preliminary studies suggest that MRI can successfully be applied to brachiure visualisation in brachiopod taxonomy, and based on the assumption that the contrast mainly stems from differences in organic matter
- ²⁵ contents, we estimate that many other types of samples will behave similarly when subjected to MR, provided that they exhibit structures where organic and anorganic are arranged in closely intertwined but separate layers – e.g. the complex septal configuations in ammonites or internal structures of fossil wood, bones and teeth. Taken together, the microscopic resolution currently achieved with Magnetic Resonance tech-

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niques, their non-invasiveness, the possibility to obtain spectroscopic and 3D spatial information, their potentially broad applicability and the multitude of ongoing efforts to further improve them (Glover and Mansfield, 2002) all suggest they could provide helpful insights into a wide range of paleobiological issues, with the precious samples remaining intact.

Supplementary information

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http://www.copernicus.org/EGU/bg/bgd/2/239/bgd-2-239-sp1.zip

Movie 1: Sublethal injury with fracture of the rostrum in *Gonioteuthis quadrata* (Blainville, 1827).

¹⁰ Movie 2: Sublethal injury of the muscle mantle in *Gonioteuthis quadrata* (Blainville, 1827).

Movie 3: Apical collar in Gonioteuthis quadrata (Blainville, 1827).

Movie 4: Disturbance of the guard secreting mantle in *Neoclavibelus subclavatus* (Voltz, 1830).

¹⁵ Movie 5: Early disturbance of the guard secreting mantle in *Hibolithes jaculoides* (Swinnerton, 1937).

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Fig. 1. Sublethal injury with fracture of the rostrum. *Gonioteuthis quadrata* (Blainville, 1827) guard with a zigzag-like deformation. **(A)**: lateral view (photomicrograph). **(B)**: median MR section. Arrow heads indicate fractures. The corresponding slice series (in steps of 125 μ m) with this orientation is given in Movie 1. Acquisition parameters: Field of view (FOV): 15×15×32 mm³. Number of averages (N_A): 2. Total acquisition time (*T_a*): 18 h. Colour scale: MR signal intensity in arbitrary units (identical for all MR images). Scale bar: 5 mm for all MR images slices.

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Fig. 2. Sublethal injury of the muscle mantle. *Gonioteuthis quadrata* (Blainville, 1827) guard with a twin apex. (A): lateral view (photomicrograph). (B): longitudinal MR section. (C): transverse MR section through the beginning of the anomaly exhibiting the preceding injury of the interimistic guard's surface. (D): transverse MR section through the initial bulg. (E): transverse MR section through the double rostrum at the posterior part. The corresponding slice series (proximal to distal in steps of 109 μ m) with transverse orientation is given in Movie 2. Acquisition parameters: FOV: $10 \times 12 \times 28 \text{ mm}^3$. N_A: 4. *T_a*: 36 h.

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Fig. 3. Apical collar. Gonioteuthis quadrata (Blainville, 1827) guard with a collar-like dorsal anomaly. **(A)**: lateral view (photomicrograph). **(B)**: median MR section. **(C)**: transverse MR section through the anomalous dorsal lip. The corresponding slice series (proximal to distal in steps of 109 μ m) with transverse orientation is given in Movie 3. FOV: 12×12×28 mm³. T_r : 852 ms. N_A: 12. T_a : 93 h.

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Fig. 4. Disturbance of the guard secreting mantle. *Neoclavibelus subclavatus* (Voltz, 1830) guard (collection of V. Kriegisch) with an anomalous protuberance, presumably caused by parasitism. **(A)**: lateral view (photomicrograph). **(B)**: median longitudinal MR section. Arrows indicate positions of the slices depicted in (C) and (D). **(C)**: transverse MR section through the bubble-like anomaly. **(D)**: more distal transverse MR section through normal tissue. The corresponding slice series (proximal to distal in steps of 117 μ m) with transverse orientation is given in Movie 4. The radial lines are indicative of the direction of calcite crystal growth. **(E)**: 3D model of the guard, directly obtained from the MR images. FOV: 8×8×30 mm³. *T_r*: 952 ms. N_A: 10. *T_a*: 87 h.

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Fig. 5. Early disturbance of the guard secreting mantle. Anomalous rostrum of *Hibolithes jaculoides* (Swinnerton, 1937). **(A)**: lateral view (photomicrograph). **(B)**: median longitudinal MR section. **(C)**: dorsal longitudinal MR section. The corresponding slice series (proximal to distal in steps of 113 μ m) with transverse orientation is given in Movie 5. FOV: 13×11×29 mm³. N_A: 4. *T_a*: 36 h.

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