

Late Cretaceous coprolite from the Opole area (southern Poland) as evidence for a variable diet in shell-crushing shark *Ptychodus* (Elasmobranchii: Ptychodontidae)

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Background. Coprolites, i. e. fossilized faeces, are an important source of knowledge on the diet and food processing mechanisms in the fossil record. Direct and indirect evidences for the dietary preferences of extinct sharks are rare in the fossil record. The first coprolite attributable to *Ptychodus* containing prey remains from the European Cretaceous is documented here.

Methods. A coprolite from the Late Cretaceous of Opole (southern Poland) was scanned using micro-computed tomography to show the arrangement of the inclusions. In addition, the cross-section was examined under the SEM/EDS to analyse the microstructure and chemical composition of the inclusions.

Results. Brachiopod shell fragments and foraminiferan shells are recognized and identified among the variously shaped inclusions detected through the performed analysis.

Conclusions. The extinct shell-crushing shark *Ptychodus* has been identified as the possible producer of the examined coprolites. The presence of brachiopod shell fragments indicates that at least some species of this durophagous predatory shark may have fed on benthic molluscs by hunting over the sea bottom.

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Abstract

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Introduction

Coprolites, i.e. fossilized faeces, together with consumulites (intestine contents), gastroliths (stomach, or gizzard, stones), and regurgitalites (orally expelled masses) make up the group of ichnofossils known as bromalites (Hunt & Lucas, 2021). These are informative for establishing the diet and food processing style. The major caveat is the uncertainty concerning the specific producer of this kind of fossils. Sometimes, the co-occurrence in the same strata of fossils and

faeces, and specific features of the animal linking the coprolite and skeletal material (e.g. size, purported diet), can be used as means to pinpoint, with a certain level of certainty, the most likely producer. This was done for the Late Triassic site of Krasiejów in the Opole area, where small coprolites, containing insect remains, were identified as a product of a co-occurring dinosauiromorph *Silesaurus opolensis*, with the main reasoning based on body sizes and possible diets of the skeletally identified fauna at this locality (Qvarnström et al. 2017, 2019, 2021). The discussion there, however, did not take into account a range of taxa from the site identified thus far only on dental remains. Shark teeth and coprolites are a common find in Late Cretaceous deposits, including the Turonian-Coniacian of Opole area (Mazurek, 2008). Skeletal fossils consist mainly of isolated teeth, with few finds of an associated dentition or even a single vertebra (pers. obs.). Niedźwiedzki (2005) and Niedźwiedzki & Kalina (2003) are the only authors that have studied the shark fauna of the Opole area in recent years. Niedźwiedzki & Kalina (2003) described from Opole the following taxa: *Ptychodus latissimus*, *P. mammillaris*, *P. polygyrus*, *Squalicorax* sp., *Scapanorhynchus raphiodon*, and *Paranomotodon angustidens*. Niedźwiedzki (2005) listed jointly taxa from localities at Opole and Sudetes area. Apart from those mentioned above, other taxa said to be common were *Cretoxyrhina mantelli*, *Cretolamna appendiculata*, *Squalicorax falcatus*, and *Odontaspis subulate*, while rare finds included *Hexanchus microdon*, *Synechodus major* and *Hybodus dentalus*. In a popular book (Yazykova (ed.) - 2022), Niedźwiedzki confirms the presence specifically in the Opole area of *Squalicorax falcatus*, *Cretolamna appendiculata*, *Cretoxyrhina mantelli*, and *Odontaspis subulata*. These works are supplemented by the collecting efforts of the current authors, whose rich collection preserves *Squalicorax falcatus* and other lamniforms, *Ptychodus* spp., as well as a single find of a hexanchiform.

As for coprolites, spiral shark faeces are especially common in clayey marls. Their general presence was already noted by Mazurek, 2008; Hunt et al. 2015). Here, we present and document in detail for the first time one of the coprolites from the Upper Cretaceous of Opole (southern Poland). The specimen was analysed by SEM-EDS and microCT to investigate structure and chemical composition of the inclusions. Based on the shape and prey content of the coprolite and the dietary preferences of the co-occurring ichthyofauna, the coprolite producer was identified and its feeding was discussed in a palaeoecological context.

behavior

Geological setting

Odra II quarry is a working quarry within the city of Opole (southern Poland). The exposed rock sequence starts with clayey marls (Middle Turonian *Inoceramus apicalis* Zone) and proceeds with limy marlstones (Middle Turonian *I. lamarcki* Zone to the lowermost part of Upper Turonian *I. perplexus* Zone), and ends with marly limestones (*I. perplexus* Zone). This sequence of strata forms part of a single transgression-regression megacycle (Cenomanian-Coniacian) that represents the Cretaceous strata of the so-called Opole Trough (Jagt-Yazykova et al. 2022). The biota preserved is numerous and consists of ichnofossils, sponges, inoceramids and other

bivalves, brachiopods, fish remains, cephalopods, echinoderms, crustaceans, cnidarians, shark coprolites, land flora, and rare marine reptiles. The coprolites are quite common and of uniform size and shape, with spiral structure pointing to sharks ~~as their makers~~. The specimen studied comes from the clayey marls (Middle Turonian: *I. apicalis* Zone).

→ Dentzien-Dias et al. 2012

Materials & Methods

A coprolite was collected from the Odra II quarry (Oleska street, Opole) during the summer digging camp in 2020. It is housed at University of Opole (col. no. IBUO-DM-KOPRO1). Fieldwork was possible due to the legal agreement between the quarry owner (Cement Factory “Odra” and European Centre of Palaeontology, University of Opole) dated 24.05.2017. The coprolite is incomplete and the preserved portion is 22 mm in length. The estimated size of the coprolite could be at least two times larger compared to other specimens in the collection ranging between ca. 20-55 mm in total length. As the specimen is broken, some dark infillings are visible within the grey phosphatic mass on the cross-section (Fig. 1). To determine the composition of the infilling, the specimen was analysed with micro CT scanner SkyScan 1273 in Bruker Laboratory in Kontich, Belgium. Obtained data were presented using DataViewer (for multiple cross sections in three directions) and CTVox (for the presentation of the 3D orientation of infillings) software. 8.5µm resolution scan is uploaded to Morphosource database (<http://n2t.net/ark:/87602/m4/514300>) in the form of 2882 tiff image series. For chemical identification of the infilling, the surface of the broken part (cross-section) was polished with grinding powder. The obtained polished surface was examined under Scanning Electron Microscope TM 3000 with secondary electrons as well as with the use of Energy-Dispersive X-ray Spectroscopy. In addition, the coprolite IBUODM-KOPRO2 (Fig. 1C) was selected as comparison material.

Results

Examined specimen and additional IBUIDM-KOPRO2 possess a heteropolar spiral shape (see also Fig. 2D), which is typical of chondrichthyan coprolites (see Eriksson et al. 2011; Dentzien-Dias et al. 2012). MicroCT scans reveals numerous infillings with densities differing from the phosphatic matrix of the coprolite (Fig. 2, 3). Most of the shapes are irregular, many being boat-shaped. Some of them can be recognized and assigned to certain groups of animals, specifically micromorphic brachiopods (Fig. 4) and foraminifera (Figure 2F), based on, SEM observations of microstructure and cross-section visible in micro CT scan. Two unidentified shells/tests have been observed under higher magnification under SEM. Both inclusions (Fig. 4) show the walls consisting of horizontal lamellae. No vertical elements are present, which would be expected in the case of an inoceramid prismatic layer (e. g. Jiménez-Berrocso et al., 2006), one of the possible prey. No macroscopic chunks of large bivalves are present either. The microstructure is more reminiscent of an inopuncate brachiopod shells (Griesshaber et al., 2007). Regardless, some inclusions are firmly identified as brachiopods and forams (Fig. 2, 3), while no traces of other possible shelled (e.g. inoceramids, see Hattin, 1975) or soft-bodied prey were detected.

In the EDS analysis, the main elements are Ca, O, C, and P (Fig. 4).

Discussion

Irregular and boat-shaped infilling creates a specific pattern. ~~Similar can~~ be observed in coprolites of durophagous fishes from the Middle Triassic (Antczak et al., 2020). EDS signature suggests that these are elements made of calcium carbonate. The matrix of the coprolite possesses a phosphatic character. The spiral heteropolar nature of the Opole Cretaceous coprolites points to sharks as their producers. Taking into account the above, it strongly suggests that the analysed coprolite was produced not by a piscivorous shark but rather by species feeding on invertebrates with calcareous shells. The only known candidate is *Ptychodus*. Currently, this genus is thought to be a facultative durophage, with diet composed of inoceramids and other shelly fauna, but also fishes (Shimada et al. 2009; Amadori et al. 2019, 2020, 2023; Hamm 2020). The assignment of some of the infillings to brachiopods suggests that the producer was feeding at the bottom of the sea (nekto-benthonic) instead of in open water (nektonic). In addition, tests of calcareous foraminifera can be recognized, similar to genera *Lenticulina* or *Gavelinella* (Kłapciński & Teisseyre, 1981) which are bottom-dwelling taxa, probably swallowed accidentally together with the sediment and a brachiopod laying on the bottom of the sea.

In the Turonian of Opole, several shark species could produce coprolites of this size. The known taxa are *Cretoxyrinha*, *Hexanchus*, *Squalicorax*, and *Ptychodus*. Among them, only the last is commonly described as durophagous based on tooth morphology (Shimada et al., 2009, 2010) (Fig. 5). Niedźwiedzki and Kalina (2003) identified at the Opole Cretaceous three taxa of *Ptychodus*. Apart from isolated teeth, the Opole Cretaceous also yielded two sets of teeth: one is deposited at the University of Wrocław, while the other is in a museum of the University of Opole. Similar finds are known for several taxa worldwide (Amadori et al., 2019; Hamm, 2017), with partial skeletons or skulls much rarer (Shimada et al., 2009, 2010).

Occurrence of *Ptychodus* as the only durophagous shark suggests that the producer of the coprolite might be specifically identified to the mentioned genus. However, the lack of shell fragments within the coprolite is notable. There are several possible explanations.

First is that producer of the coprolite fed also on the common inoceramids, but was able to feed only on the soft tissue and for example orally reject the hard shells. The modern mammal *Odobenus rosmarus* feeds on benthic mollusks by sucking the soft tissue and ejecting the hard parts (Sheyer et al., 2011). However, currently, no dentalites were recognized from Opole Cretaceous inoceramid shells (even though many microscopical epifauna remnants can be observed – e. g. Bryozoa, Serpulidae, Ostreoida). From numerous specimens described by Walaszczyk (1992) a single sublethal injury was mentioned. If sharks were efficient predators we would predict evidence of failed prey subjugation. However deformations and growth iterations in inoceramid shells are known, they are rather effects of decapod predation (Harries & Ozanne, 1998). Of note, none of the coprolites we studied externally seem to contain any large shelly material. To the best of our knowledge not such are known elsewhere.

The second possibility is that the fossils of a coprolite producer are not present (or not recognized yet) in the Quarry due to the sedimentation bias or being less common representative

Similar infillings

collection numbers?

of the Cretaceous fauna of this area. Hunt et al. (2015) show that producers of coprolites are often not represented by body fossils. Chondrichthyan fossilized faeces are the most common, while in terms of body fossils palaeoichthyofaunas are usually much more diversified, which Hunt et al. (2015) termed the 'shark surplus paradox'. The third option, explaining this the lack of dentalites and brachiopod infillings in the described coprolite is to consider *Ptychodus* (the form from Opole, and by extensions possibly also other members of the genus), as the producer which, contrary to some current opinions, was not a strictly durophagous taxon, but rather a durophagous-filter feeder specialized in small prey, with bulbous teeth for crushing shells, but also with water moving between the ridges of the teeth (Fig. 1). Such elaborated ornamentation as present on the teeth of *Ptychodus* is lacking in many other durophagous taxa except skates, including among others: fishes (e. g. Purnell and Darras, 2015; Raguin et al., 2020), placodonts (Pommery et al., 2021) and many mosasaurs (Leblanc et al., 2019), the teeth are often restricted to the outer edge of the jaws, and supposed shark dentalites on inoceramids and other hard elements are rare in the literature known to us (e.g. Kauffman, 1972; Hunt & Lucas, 2021, table A.5), which however can be ascribed to poor taphonomic potential of such finds, and lack of both recognition and studies devoted to them. Also not all filter-feeders possess small, gracile, sieve-like teeth. Several species of pinnipeds have teeth modified into filter-feeding, specifically with elaborate cusps of postcanines on both the upper and lower jaw. This modification is well-seen, especially in the crabeater seal *Carinophaga lobodon* (Chatterjee & Small, 1989; Bengtson, 2002; Adam, 2005).

*Based
on
fishes*

Conclusions

MicroCT scan and EDS analysis show that coprolite collected in the Turonian deposits of Odra II quarry in Opole, southern Poland is filled with shell fragments. Inclusions can be identified as remains of small brachiopods (and occasionally Foraminifera). Such content suggest that the producer's diet was based on the small shell-covered organisms living within the sediment, possibly revealing mix of durophagy and filter-feeding strategy. According to the shape of the coprolite it can be described as belonging to shark. Within chondrichthyan fauna of the locality there is only one species of durophagous shark, *Ptychodus*, thus it can be proposed as the producer of the analysed coprolite, although 'shark surplus paradox' need to be considered as well (Hunt et al., 2015). *Ptychodus* (if considered a producer) might have been a durophagous-filter feeder and not a strictly durophagous fish as there is no evidence of preying on abundant large inoceramids and other shelly organisms (in the forms of coprolites or regurgitalites). While we acknowledge this hypothesis don't necessary be universally applied to other species of the genus, or different growth stages – in the context of scarcity of direct evidence worldwide for preying on large shelly organisms, we tentatively suggest that some form of both durophagy and filter feeding ecology might need to be considered for *Ptychodus* spp. individuals. Further investigation of

coprolites and, when available, gut contents will be necessary to confirm or reject the hypotheses proposed in this study.

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324 Fig. 1. *Ptychodus* remains from Opole Cretaceous. Analysed coprolite IBUO-DM-KOPRO1 in
325 lateral view (A) and cross-section (B). Coprolite IBUO-DM-KOPRO2 in lateral view (C). Teeth
326 IBUO-DM-ZAB1 (D).

327 Fig. 2. MicroCT scan of the coprolite. Infillings – 3D model (A-B). Coprolite mass with
328 infillings – 3D model (C-D). Longitudinal cross-section (E-F). b – brachiopod shell, f – foram
329 shell. S – spiral structure.

330 *Gavelienella* illustration from Hornibrook et al. 1989, Fig. 18.17. Brachiopod shell photograph
331 from alexstrekeisen.it. 3D model made in CTVox. Scan resolution: 8.5µm

332 Fig. 3. Cross-sections of the analysed coprolite in 3 directions (A, C, D). Magnification of the
333 example of indet. shell fragment (B). Image obtained in DataViewer

334 Fig. 4. EDS analysis. Brachiopod shell fragments (A, B), the surface of the EDS analysis (C),
335 and mass percentage result (D). SEM photographs: own. Made at Faculty of Chemistry,
336 University of Opole.

337 Fig. 5. *Ptychodus* reconstruction(Author: Jakub Kowalski) with an example of tooth IBUO-DM-
338 ZAB1 and coprolite (IBUO-DM-KOPRO2).

339

340

Figure 1

Ptychodus remains from Opole Cretaceous.

Analysed coprolite IBUO-DM-KOPRO1 in lateral view (A) and cross-section (B). Coprolite IBUO-DM-KOPRO2 in lateral view (C). Teeth IBUO-DM-ZAB1 (D).

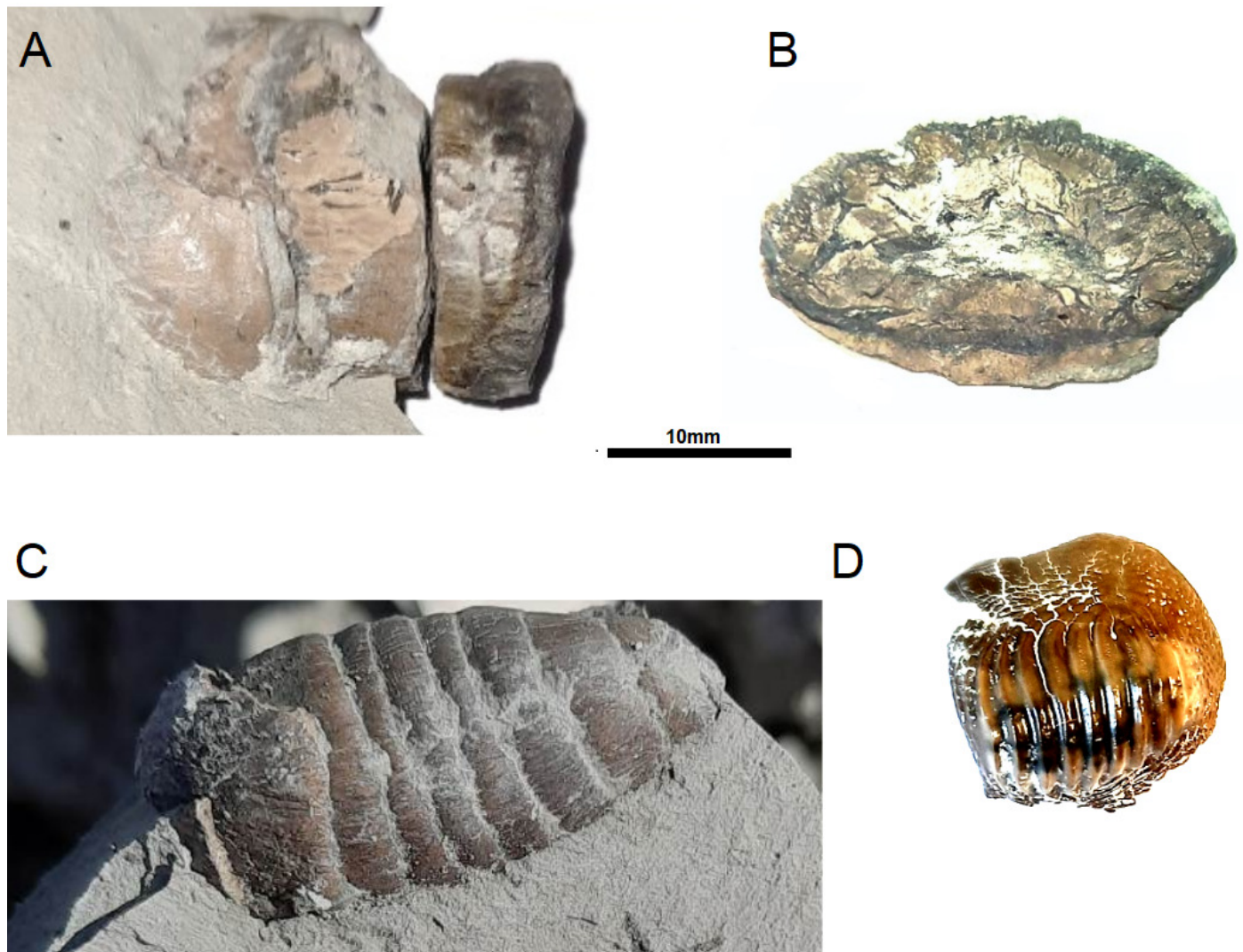


Figure 2

MicroCT scan of the coprolite.

Infillings - 3D model (A-B). Coprolite mass with infillings - 3D model (C-D). Longitudinal cross-section (E-F). b - brachiopod shell, f - foram shell. S - spiral structure. *Gavelienella* illustration from Hornibrook et al. 1989, Fig. 18.17. Brachiopod shell photograph from alexstrekeisen.it. 3D model made in CTVox. Scan resolution: 8.5µm

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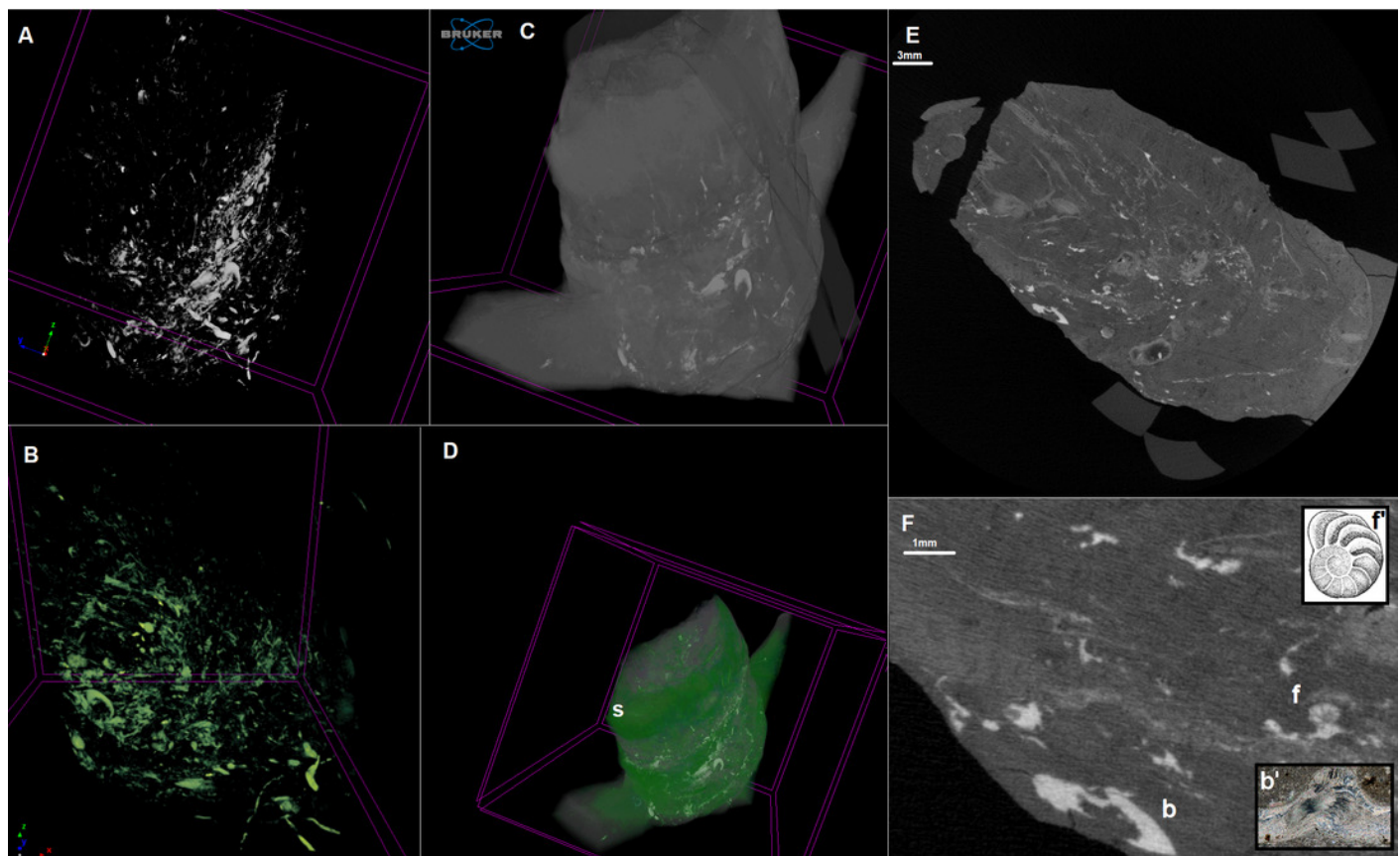


Figure 3

Cross sections of the analysed coprolite.

Cross-sections of the analysed coprolite in 3 directions (A, C, D). Magnification of the example of indet. shell fragment (B). Image obtained in DataViewer.

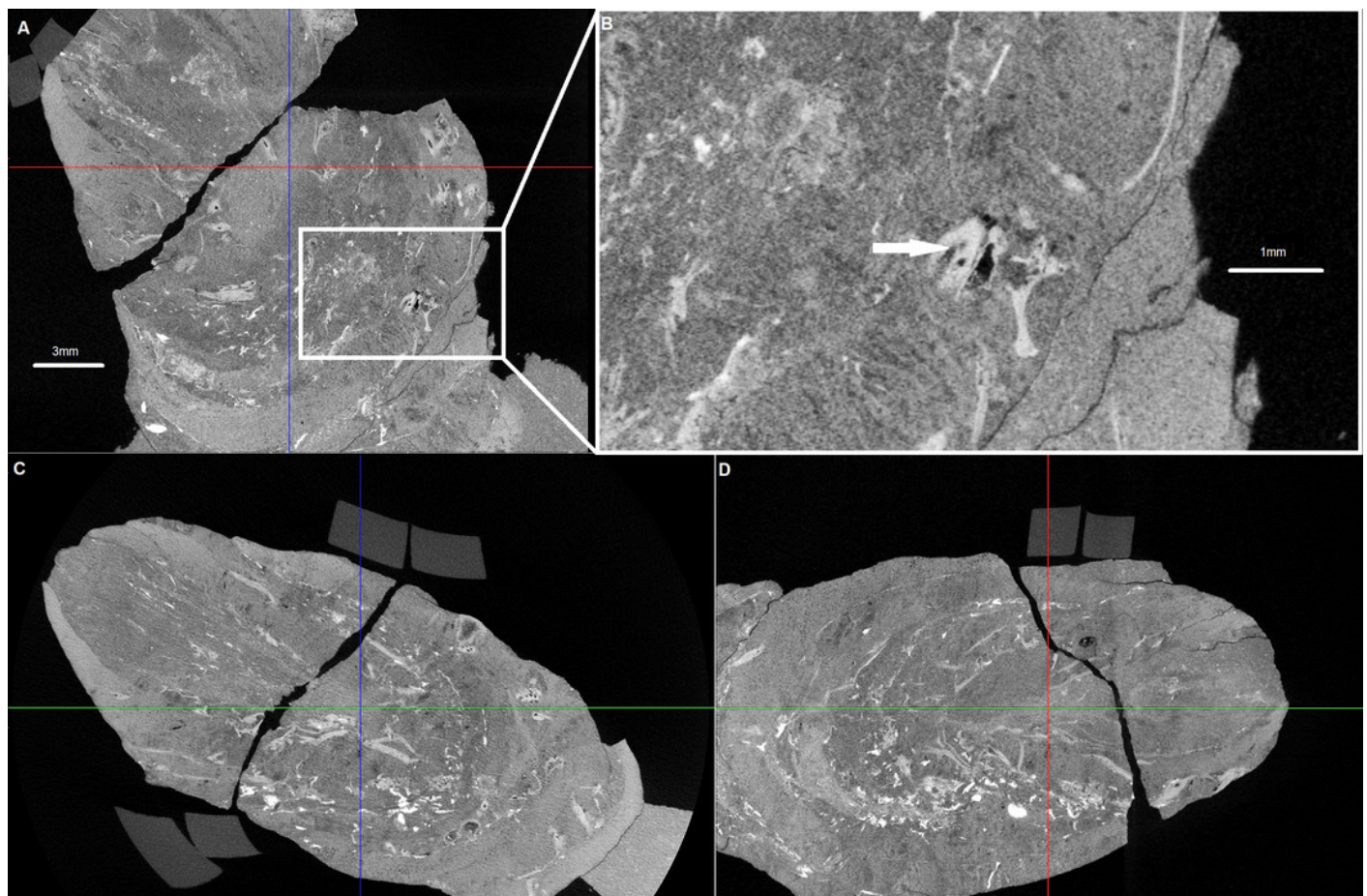


Figure 4

Fig. 4. EDS analysis.

Brachiopod shell fragments (A, B), the surface of the EDS analysis (C), and mass percentage result (D). SEM photographs: own. Made at Faculty of Chemistry, University of Opole.

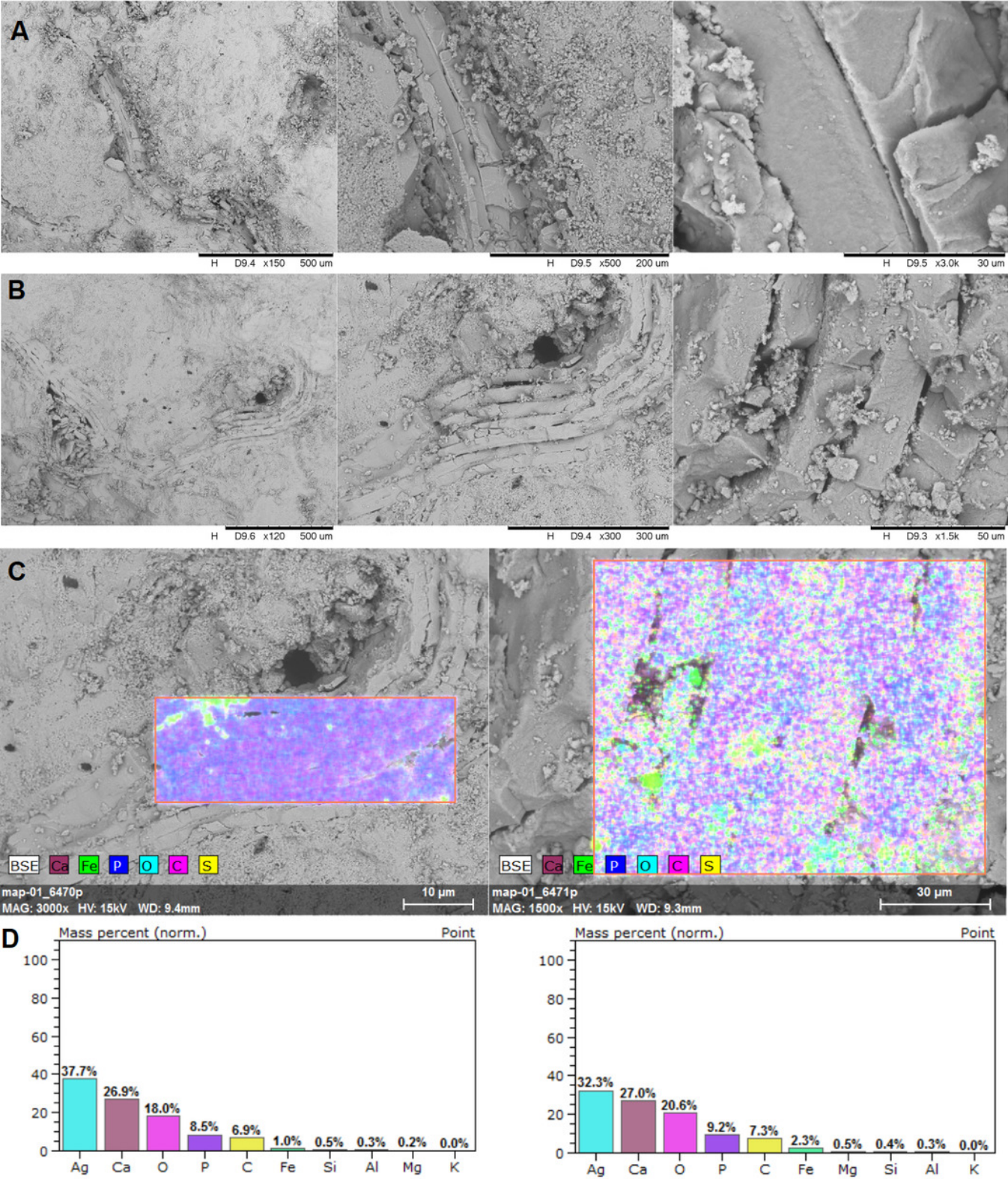


Figure 5

Ptychodus reconstruction (Author: Jakub Kowalski) with an example of tooth IBUO-DM-ZAB1 and coprolite (IBUO-DM-KOPRO2).

