



Rahman, I., Waters, J. A., Sumrall, C. D., & Astolfo, A. (2015). Early postmetamorphic, Carboniferous blastoid reveals the evolution and development of the digestive system in echinoderms. Biology Letters, 11(10), [20150776]. DOI: 10.1098/rsbl.2015.0776

Peer reviewed version

License (if available): Unspecified

Link to published version (if available): 10.1098/rsbl.2015.0776

Link to publication record in Explore Bristol Research PDF-document

This is the author accepted manuscript (AAM). The final published version (version of record) is available online via The Royal Society at DOI: 10.1098/rsbl.2015.0776. Please refer to any applicable terms of use of the publisher.

University of Bristol - Explore Bristol Research General rights

This document is made available in accordance with publisher policies. Please cite only the published version using the reference above. Full terms of use are available: http://www.bristol.ac.uk/pure/about/ebr-terms.html

1	Early post-metamorphic, Carboniferous blastoid reveals
2	the evolution and development of the digestive system
3	in echinoderms
4	
5	Imran A. Rahman ^{1*} , Johnny A. Waters ² , Colin D. Sumrall ³ and Alberto Astolfo ⁴
6	
7	¹ School of Earth Sciences, University of Bristol, Life Sciences Building, 24 Tyndall Avenue,
8	Bristol BS8 1TQ, UK.
9	² Department of Geology, Appalachian State University, Boone, NC 28608, USA.
10	³ Department of Earth and Planetary Sciences, University of Tennessee, Knoxville, TN 37996,
11	USA.
12	⁴ Swiss Light Source, Paul Scherrer Institut, Villigen 5232, Switzerland.
13	
14	*Author for correspondence:
15	Imran A. Rahman

16 e-mail: <u>imran.rahman@bristol.ac.uk</u>

17 Abstract

Inferring the development of the earliest echinoderms is critical to uncovering the 18 evolutionary assembly of the phylum-level body plan but has long proven problematic 19 20 because early ontogenetic stages are rarely preserved as fossils. Here, we use synchrotron tomography to describe a new early post-metamorphic blastoid echinoderm from the 21 Carboniferous (~323 Ma) of China. The resulting three-dimensional reconstruction reveals a 22 U-shaped tubular structure in the fossil interior, which is interpreted as the digestive tract. 23 Comparisons with the developing gut of modern crinoids demonstrate that crinoids are an 24 25 imperfect analogue for many extinct groups. Furthermore, consideration of our findings in a phylogenetic context allows us to reconstruct the evolution and development of the digestive 26 system in echinoderms more broadly; there was a transition from a straight to a simple curved 27 28 gut early in the phylum's evolution, but additional loops and coils of the digestive tract (as seen in crinoids) were not acquired until much later. 29

30

```
31 Keywords:
```

32 echinoderms, blastoids, evolution, development, synchrotron tomography

33

34

35 1. Background

36 Reconstructing the origin and evolution of animal body plans requires a detailed

understanding of the developmental processes underpinning changes in adult morphology [1],

and echinoderms are a model group in this respect. Select species have been a focus of

39 investigation for over a century, and recent work on developmental gene regulatory networks

40 has revolutionized our understanding of cell specification mechanisms [2]. Most scenarios for

41 the evolution of development in echinoderms are thus founded chiefly on extant taxa [3].

42 However, living echinoderms have diverged greatly from the latest common ancestor they share with all deuterostomes and, hence, might not closely reflect developmental processes in 43 the earliest forms. Study of the extensive Palaeozoic fossil record of echinoderms could shed 44 light on the early evolution of development in the phylum, but such work is hampered by the 45 scarcity of fossilized early ontogenetic stages (although see [4]). Here, we report an 46 extremely well-preserved, early post-metamorphic echinoderm from the Carboniferous (~323 47 48 Ma) of China. The fossil belongs to Blastoidea, an extinct clade of stemmed echinoderms (pelmatozoans) with pentaradial symmetry [5]. Synchrotron tomography was used to study 49 50 the internal anatomy of this specimen, thereby revealing, for the first time, preserved evidence of the digestive tract in a blastoid echinoderm. This informs on the development of 51 the blastoid gut, with implications for elucidating the evolution and development of the 52 53 digestive system in early echinoderms in general.

54

55

56 2. Material and methods

The studied specimen (NHMUK EE 15671) comes from the Mississippian (lower 57 58 Carboniferous) Luocheng Formation, Xinxu, Guangxi, China [6], and is housed in the Natural History Museum, London. Propagation-based phase-contrast synchrotron radiation 59 X-ray tomographic microscopy was performed at the TOMCAT beamline of the Swiss Light 60 Source, Paul Scherrer Institut, Villigen, Switzerland. The fossil was scanned using an X-ray 61 energy of 20 keV, 1501 projections, and an exposure time of 200 ms, and the sample-to-62 detector propagation distance was set at 40 mm. This gave a tomographic dataset with a voxel 63 size of 0.65 µm (electronic supplementary material, data S1), which was digitally 64 reconstructed as a three-dimensional virtual fossil (electronic supplementary material, data 65 66 S2, video S1) using the SPIERS software suite [7].

67

68

69 **3. Results**

70 The specimen measures 2.85 mm in height and 1.1 mm in maximum width (figure 1). The theca is cone-shaped and consists of three well-developed basal plates and five well-71 72 developed radial plates (figure 1a,d). The mouth is centrally located at the uppermost extremity of the theca (the summit) and is bordered by the deltoid plates, which are small and 73 represented by the deltoid lips (figure 1a,b). The anus, proximal edges of the spiracles, and 74 75 terminal food grooves are also situated on the summit. Each of the food grooves leads to three brachiole facets and is presumably supported by an internal lancet plate. Adjacent to the 76 anus, a notch in the CD interray (in the C and D ray radials) marks the future position of the 77 78 hypodeltoid plate, and a small wedge of calcite in this notch might reflect the nucleation of the plate (figure 1*a*,*b*). Medial notches in the radials likely represent the initiation of the 79 80 ambulacral sinuses. Each of the basals and radials is covered by distinct external growth lines (figure 1*a*,*b*,*d*). 81

The presence of three basals, five radials, and the configuration of the summit 82 83 unequivocally demonstrate that the specimen is a blastoid. The small size of the theca and the absence of the anal deltoids, the deltoid bodies, and the hydrospires indicate that the 84 specimen was immature; the presence of the deltoid lips, the most proximal elements of the 85 86 ambulacral feeding system, and (possibly) the lancets strongly suggests that the specimen had progressed past the passalocrinid stage of blastoid development (as defined by [8]). The 87 passalocrinid stage was inferred to be equivalent to the non-feeding cystidean stage of crinoid 88 89 ontogeny [8]; the studied specimen represents a more advanced post-metamorphic developmental stage that was capable of feeding (as shown by the presence of brachiole 90 91 facets associated with the food grooves), analogous to the pentacrinoid stage of crinoid

92 ontogeny. The fossil cannot be identified to the genus or species level because it lacks a93 number of key adult characters.

The basals and radials are characterized by a core of microperforate stereom, in which 94 the pores are small and circular in cross section (up to about 15 µm in diameter), and 95 frequently filled with pyrite (figure 1f). The thecal plates surround a large cavity, oval to 96 97 pentagonal in cross section, which is occupied by fine-grained sediment (partially pyritized) and connects to the mouth and anus at the summit. A small U-shaped tubular structure is 98 situated within the upper portion of the cavity (figure 1c,e,f). This tube is circular to elliptical 99 100 in cross section, measuring approximately 80–190 µm in diameter and approximately 2.5 mm in total length. It consists of a thin rim (approximately 5–8 µm in width) surrounding a fill of 101 102 fine-grained sediment (figure 1*f*). The tube follows a gently undulating course from one end 103 to the other, with occasional sharp kinks, and it decreases in diameter noticeably at the proximal end. Both 'arms' of the 'U' extend towards the summit where the mouth and anus 104 are located; however, no direct connection to these orifices is preserved (figure 1c,e). 105

106

107

108 4. Discussion

The U-shaped tubular structure described herein (figure 1c,e,f) has not previously been 109 reported in a fossil blastoid. One possibility is that the tubular structure represents a vertical 110 U-shaped trace fossil; however, there is no evidence of external boring and the U-shaped tube 111 is not connected to the exterior of the specimen (figure 1c,e), strongly arguing against this 112 interpretation. Instead, the morphology and position of the tubular structure are more 113 consistent with what is described for the digestive tract of extant crinoids [9–11]; in both 114 taxa, the tube is broadly U-shaped and restricted to the central region of the body. 115 116 Consequently, we interpret the tubular structure in the blastoid fossil as an incompletely

preserved part of the digestive system, hitherto unknown in blastoids, and it is inferred that the two upwardly projecting 'arms' of the 'U' were connected to the mouth and anus in life, as in the pentacrinoid stage of crinoid development [11].

120 Crinoids are the only extant members of the pelmatozoan clade, which is represented predominantly by a number of extinct Palaeozoic groups (e.g. blastoids, eocrinoids, and 121 rhombiferans). As a result, crinoids are often assumed to be a close analogue for the anatomy 122 123 and development of fossil pelmatozoans, including blastoids [5], and they have even been taken as an exemplar for basal echinoderms more generally [3]. However, comparisons with 124 125 the post-metamorphic blastoid described in this study demonstrate that crinoids are unlikely to be a good interpretive model for many fossil groups. In the comatulid crinoid Antedon, the 126 digestive tract forms soon after metamorphosis, arising from the enteric sac in the cystidean 127 128 stage as a looped tubular structure that coils in a clockwise direction. By the pentacrinoid stage, a connection to the anus has been established, and the digestive tract becomes 129 differentiated into oesophagus, stomach, and intestine before the onset of the free-living 130 juvenile stage [9–11]. The post-metamorphic ontogeny of stalked crinoids is less well known, 131 but recent work on the isocrinid *Metacrinus rotundus* is suggestive of a similar 132 developmental history, with the digestive tract (mouth and oesophagus) forming in the 133 cystidean stage [12]. Thus, the developing gut of crinoids differs from what we describe for 134 135 blastoids, where it takes the form of a simple U-shaped curve, with no evidence of looping, 136 clockwise coiling, or differentiation into distinct organs (figure 1c,e). Whereas it is possible that differentiation of the digestive tract occurred later in blastoid ontogeny, further structural 137 changes, such as looping and coiling, can be considered highly improbable because these are 138 139 established before the feeding pentacrinoid stage in modern crinoids [11]. This strongly suggests that the development of extant crinoids involved more radical changes to the 140

6

organization of the internal organs, including the digestive system, than occurred in blastoidontogeny, and the same may be true for early pelmatozoans in general.

There is very little information about the morphology of the gut in most groups of 143 fossil echinoderms, but the general structure can be inferred from the position of the mouth 144 and anus (figure 2). The most basal stem-group echinoderms (Ctenoimbricata and 145 ctenocystoids) have an anterior mouth and a posterior anus [13], suggesting a more or less 146 147 straight gut. However, in more crownward stem-group taxa (cinctans and solutes), the anus is typically situated closer to the mouth, indicating that the gut had become broadly U-shaped or 148 149 J-shaped. This shape was likely established early during development, i.e. metamorphosis, as there is no evidence of radical morphological changes during later juvenile development [14]. 150 151 Our description of the digestive system in a post-metamorphic blastoid suggests that a simple 152 curved gut was also present in basal pelmatozoans; the absence of additional developmental 153 transformations of the gut (i.e. looping or clockwise coiling), such as those seen in crinoids (including fossil forms, see [15]), indicates that these changes are most probably a derived 154 character of some pelmatozoan groups (perhaps restricted to crinoids). This contradicts the 155 hypothesis that the pentaradial organization of echinoderms evolved as a consequence of the 156 formation of multiple loops of the digestive tract [16], as well as the theory that the 157 placement of the anus in the CD interray was the product of clockwise coiling of the gut [17]. 158 Instead, the results presented herein are more compatible with [18], who hypothesized the 159 160 presence of a simple U-shaped gut in early echinoderms.

161

162

163 Data accessibility. The original slice images, the digital reconstruction, and a video file are
164 available at: http://dx.doi.org/10.5061/dryad.sq134.

165

166	Author contributions. I.A.R., J.A.W., and C.D.S. designed the project. I.A.R., J.A.W., and					
167	A.A. carried out scans. I.A.R. produced the digital reconstruction. I.A.R. wrote the first draft					
168	of the manuscript and all authors contributed to subsequent versions. All authors gave final					
169	approval for publication.					
170						
171	Competing interests. We have no competing interests.					
172						
173	Funding. I.A.R. was funded by an 1851 Royal Commission Research Fellowship. J.A.W.					
174	and C.D.S. were supported by grants from the National Science Foundation (DEB-1036416					
175	and DEB-1314236).					
176						
177	Acknowledgements. We are grateful to Sara Marcus (Oregon Institute of Technology) for					
178	collecting the specimen and acknowledge the Paul Scherrer Institut, Villigen, Switzerland for					
179	the provision of synchrotron radiation beamtime on the TOMCAT beamline at the Swiss					
180	Light Source.					
181						
182						
183	References					
184	1. Erwin DH, Laflamme M, Tweedt SM, Sperling EA, Pisani D, Peterson KJ. 2011 The					
185	Cambrian conundrum: early divergence and later ecological success in the early history of					
186	animals. <i>Science</i> 334 , 1091–1097.					
187	2. McClay DR. 2011 Evolutionary crossroads in developmental biology: sea urchins.					
188	Development 138 , 2639–2648.					
189	3. David B, Mooi R. 2014 How Hox genes can shed light on the place of echinoderms					

among the deuterostomes. *Evodevo* **5**, 22.

- 4. Kammer TW, Bartels C, Ausich WI. In press Presumed postlarval pentacrinoids from the
 Lower Devonian Hunsruck Slate, Germany. *Lethaia*.
- 193 5. Breimer A, Macurda DB, Jr. 1972 The phylogeny of the fissiculate blastoids. *Verh. Kon.*
- 194 *Ned. Akad. Wetensch., Afd. Natuurk.* **26**, 1–390.
- 195 6. Waters JA, Webster GD. 2012 The paleogeography of Pennsylvanian crinoids and
- 196 blastoids. In Earth and Life. Global Biodiversity, Extinction Intervals and Biogeographic
- 197 *Perturbations through Time* (ed. JA Talent), p. 831–847. London, UK: Springer.
- 198 7. Sutton MD, Garwood RJ, Siveter DJ, Siveter DJ. 2012 SPIERS and VAXML; a software
- 199 toolkit for tomographic visualisation and a format for virtual specimen interchange.
- 200 *Paleontol. Electron.* **15/5T,** 14 pp.
- 8. Sevastopulo GD. 2005 The early ontogeny of blastoids. *Geol. J.* **40**, 351–362.
- 202 9. Seeliger O. 1982 Studien zur Entwicklungsgeschichte der Crinoiden (*Antedon rosacea*).
- 203 *Zool. Jahrb. Abt. Anat. Ontog. Tiere* **6**, 161–444.
- 204 10. Hyman LH. 1955 The Invertebrates: Echinodermata. New York, NY: McGraw-Hill.
- 11. Engle S. 2013 Ultrastructure and Development of the Body Cavities in Antedon bifida
- 206 (Pennant, 1777) (Comatulida, Crinoidea). http://edocs.fu-
- berlin.de/diss/receive/FUDISS_thesis_000000040355. Unpublished PhD thesis.
- 208 12. Amemiya S, Omori A, Tsurugaya T, Hibino T, Yamaguchi M, Kuraishi R, Kiyomoto M,
- 209 Minokawa T. In press Early stalked stages in ontogeny of the living isocrinid sea lily
- 210 *Metacrinus rotundus. Acta. Zool.*
- 211 13. Zamora S, Rahman IA, Smith AB. 2012 Plated Cambrian bilaterians reveal the earliest
 212 stages of echinoderm evolution. *PLoS ONE* 7, e38296.
- 213 14. Zamora S, Rahman IA, Smith AB. 2013 The ontogeny of cinctans (stem-group
- Echinodermata) as revealed by a new genus, *Graciacystis*, from the middle Cambrian of
- 215 Spain. *Palaeontology* **56**, 399–410.

216	15. Haugh BN. 1975 Digestive and coelomic systems of Mississippian camerate crinoi				
217	Paleontol. 49 , 472–493.				

- 16. Gudo M. 2005 Körperkonstruktion und evolutionäre Trends fossiler Echinodermen
 (Homalozoa, Blastoidea, Edrioasteroidea). *Senck. Leth.* 85, 39–62.
- 220 17. Rozhnov SV. 2012 The anteroposterior axis in echinoderms and displacement of the
- 221 mouth in their phylogeny and ontogeny. *Biology Bull.* **39**, 162–171.
- 18. Peterson KJ, Arenas-Mena C, Davidson EH. 2000 The A/P axis in echinoderm ontogeny
 and evolution: evidence from fossils and molecules. *Evol. Dev.* 2, 93–101.
- 224
- 225

226 Figure legends

- **Figure 1.** Early post-metamorphic, Carboniferous blastoid (NHMUK EE 15671). (*a*,*b*,*d*)
- 228 Scanning electron microscope images. (c,e) Digital reconstructions. (f) Slice image. (a)
- Angled upper view. (b) Angled upper view, plates given false colour. (c) Angled upper view,
- theca partly transparent. (d) Lateral view. (e) Lateral view, theca partly transparent. (f)
- 231 Transverse section. Abbreviations: A–E, ambulacra; an, anus; bf, brachiole facets; bp, basal
- 232 plate; dp, deltoid plates; fg, food groove; gl, growth lines; hd, hypodeltoid; mo, mouth; rp,
- radial plate; sp, stereom pores; ts, tubular structure. Colours: cyan, radial plate; green,
- brachiole facets; red, deltoid plates. Scale bars: 0.5 mm.
- 235
- Figure 2. Phylogeny of extant and extinct echinoderms and hemichordates with

reconstructions of the digestive systems (shown in blue). Abbreviations: an, anus; mo, mouth.

238

239

240 Supplementary material

241	Data S1. Synchrotron	tomography scan of	of the early post-m	netamorphic, C	arboniferous
-----	----------------------	--------------------	---------------------	----------------	--------------

blastoid (NHMUK EE 15671). The slice images are in TIFF format, compressed in a ZIParchive.

244

- 245 Data S2. Interactive three-dimensional digital reconstruction of the early post-metamorphic,
- 246 Carboniferous blastoid (NHMUK EE 15671). The reconstruction is in VAXML format,

247 compressed in a ZIP archive, and has been downsampled to reduce triangle count. To view:

248 unpack the .zip file, install the SPIERS software suite (program and documentation available

- from www.spiers-software.org), and double-click the unpacked .vaxml file.
- 250
- 251 Video S1. Video showing a three-dimensional digital reconstruction of the early post-
- 252 metamorphic, Carboniferous blastoid (NHMUK EE 15671).









