

Philip Conrad James Donoghue

List of Publications by Year in descending order

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217
papers

16,690
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16451
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h-index

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docs citations

228
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12951
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#	ARTICLE	IF	CITATIONS
1	The Ediacaran origin of Ecdysozoa: integrating fossil and phylogenomic data. <i>Journal of the Geological Society</i> , 2022, 179, .	2.1	21
2	Integrated phylogenomics and fossil data illuminate the evolution of beetles. <i>Royal Society Open Science</i> , 2022, 9, 211771.	2.4	117
3	Dietary inference from dental topographic analysis of feeding tools in diverse animals. <i>Methods in Ecology and Evolution</i> , 2022, 13, 1464-1474.	5.2	1
4	A species-level timeline of mammal evolution integrating phylogenomic data. <i>Nature</i> , 2022, 602, 263-267.	27.8	84
5	Eukaryogenesis and oxygen in Earth history. <i>Nature Ecology and Evolution</i> , 2022, 6, 520-532.	7.8	48
6	Increasing morphological disparity and decreasing optimality for jaw speed and strength during the radiation of jawed vertebrates. <i>Science Advances</i> , 2022, 8, eabl3644.	10.3	16
7	Functional assessment of morphological homoplasy in stem-gnathostomes. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2021, 288, 20202719.	2.6	8
8	Coevolution of enamel, ganoin, enameloid, and their matrix SCPP genes in osteichthyans. <i>IScience</i> , 2021, 24, 102023.	4.1	27
9	Fossil data support a pre-Cretaceous origin of flowering plants. <i>Nature Ecology and Evolution</i> , 2021, 5, 449-457.	7.8	59
10	Experimental taphonomy of organelles and the fossil record of early eukaryote evolution. <i>Science Advances</i> , 2021, 7, .	10.3	21
11	Exceptionally preserved early Cambrian bilaterian developmental stages from Mongolia. <i>Nature Communications</i> , 2021, 12, 1037.	12.8	10
12	Acanthodian dental development and the origin of gnathostome dentitions. <i>Nature Ecology and Evolution</i> , 2021, 5, 919-926.	7.8	14
13	Empirical distributions of homoplasy in morphological data. <i>Palaeontology</i> , 2021, 64, 505-518.	2.2	9
14	Diversification dynamics of total-, stem-, and crown-groups are compatible with molecular clock estimates of divergence times. <i>Science Advances</i> , 2021, 7, .	10.3	7
15	The developmental biology of <i>< i>Charnia</i></i> and the eumetazoan affinity of the Ediacaran rangeomorphs. <i>Science Advances</i> , 2021, 7, .	10.3	36
16	Fossilization processes have little impact on tipâ€calibrated divergence time analyses. <i>Palaeontology</i> , 2021, 64, 687-697.	2.2	4
17	Phylogenetic sampling affects evolutionary patterns of morphological disparity. <i>Palaeontology</i> , 2021, 64, 765-787.	2.2	6
18	X-ray nanotomography and electron backscatter diffraction demonstrate the crystalline, heterogeneous and impermeable nature of conodont white matter. <i>Royal Society Open Science</i> , 2021, 8, 202013.	2.4	5

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19	The evolution of insect biodiversity. <i>Current Biology</i> , 2021, 31, R1299-R1311.	3.9	39
20	The evolutionary emergence of land plants. <i>Current Biology</i> , 2021, 31, R1281-R1298.	3.9	67
21	Ultrastructure and in-situ chemical characterization of intracellular granules of embryo-like fossils from the early Ediacaran Weng'an biota. <i>Palaontologische Zeitschrift</i> , 2021, 95, 611-621.	1.6	3
22	The Effect of Fossil Sampling on the Estimation of Divergence Times with the Fossilized Birth-Death Process. <i>Systematic Biology</i> , 2020, 69, 124-138.	5.6	30
23	The impact of fossil stratigraphic ranges on tip calibration, and the accuracy and precision of divergence time estimates. <i>Palaeontology</i> , 2020, 63, 67-83.	2.2	25
24	Computational Fluid Dynamics Suggests Ecological Diversification among Stem-Gnathostomes. <i>Current Biology</i> , 2020, 30, 4808-4813.e3.	3.9	13
25	Performance of A Priori and A Posteriori Calibration Strategies in Divergence Time Estimation. <i>Genome Biology and Evolution</i> , 2020, 12, 1087-1098.	2.5	9
26	Mitochondrial genomes illuminate the evolutionary history of the Western honey bee (<i>Apis mellifera</i>). <i>Scientific Reports</i> , 2020, 10, 14515.	3.3	32
27	Categorical versus geometric morphometric approaches to characterizing the evolution of morphological disparity in Osteostraci (Vertebrata, stem Gnathostomata). <i>Palaeontology</i> , 2020, 63, 717-732.	2.2	10
28	Fossil cells. <i>Current Biology</i> , 2020, 30, R485-R490.	3.9	11
29	Developmental biology of <i>< i>Helicoforamina</i></i> reveals holozoan affinity, cryptic diversity, and adaptation to heterogeneous environments in the early Ediacaran Weng'an biota (Doushantuo) Tj ETQq1 1 0.784314 rgBT /Overl...	1.0	1
30	Nucleus preservation in early Ediacaran Weng'an embryo-like fossils, experimental taphonomy of nuclei and implications for reading the eukaryote fossil record. <i>Interface Focus</i> , 2020, 10, 20200015.	3.0	15
31	The origin and rise of complex life: progress requires interdisciplinary integration and hypothesis testing. <i>Interface Focus</i> , 2020, 10, 20200024.	3.0	13
32	Data curation and modeling of compositional heterogeneity in insect phylogenomics: A case study of the phylogeny of Dytiscoidea (Coleoptera: Adephaga). <i>Molecular Phylogenetics and Evolution</i> , 2020, 147, 106782.	2.7	23
33	Disparities in the analysis of morphological disparity. <i>Biology Letters</i> , 2020, 16, 20200199.	2.3	60
34	Reprint of: "Gondolellid multielement conodont apparatus (Nicoraella) from the Middle Triassic of Yunnan Province, southwestern China". <i>Palaeogeography, Palaeoclimatology, Palaeoecology</i> , 2020, 549, 109670.	2.3	0
35	Plant Evolution: Assembling Land Plants. <i>Current Biology</i> , 2020, 30, R81-R83.	3.9	21
36	Integrated phylogenomic and fossil evidence of stick and leaf insects (Phasmatodea) reveal a Permian-Triassic co-origination with insectivores. <i>Royal Society Open Science</i> , 2020, 7, 201689.	2.4	25

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37	<p>Fleas are parasitic scorpionflies</p>. <i>Palaeoentomology</i> , 2020, 3, 641-653.	1.0	17
38	Probabilistic methods outperform parsimony in the phylogenetic analysis of data simulated without a probabilistic model. <i>Palaeontology</i> , 2019, 62, 1-17.	2.2	44
39	Middle Triassic conodont apparatus architecture revealed by synchrotron X-ray microtomography. <i>Palaeoworld</i> , 2019, 28, 429-440.	1.1	12
40	Gondolelloid multielement conodont apparatus (<i>Nicoraella</i>) from the Middle Triassic of Yunnan Province, southwestern China. <i>Palaeogeography, Palaeoclimatology, Palaeoecology</i> , 2019, 522, 98-110.	2.3	18
41	Evolution: The Flowering of Land Plant Evolution. <i>Current Biology</i> , 2019, 29, R753-R756.	3.9	12
42	Origin of horsetails and the role of whole-genome duplication in plant macroevolution. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2019, 286, 20191662.	2.6	17
43	Cellular preservation of excysting developmental stages of new eukaryotes from the early Ediacaran Wengâ€™an Biota. <i>Palaeoworld</i> , 2019, 28, 461-468.	1.1	4
44	The dermal skeleton of the jawless vertebrate <i>Tremataspis mammillata</i> (Osteostraci,) Tj ETQq0 0 0 rgBT /Overlock 1.2 Tf 50 462 Td (stem		
45	Apparatus architecture of the conodont <i>Nicoraella kockeli</i> (Gondolelloidea, Prioniodinina) constrains functional interpretations. <i>Palaeontology</i> , 2019, 62, 823-835.	2.2	4
46	The apparatus composition and architecture of <i>Erismodus quadridactylus</i> and the implications for element homology in prioniodinid conodonts. <i>Papers in Palaeontology</i> , 2019, 5, 657-677.	1.5	3
47	The circulatory system of <i>Galeaspida</i> (Vertebrata; stem-Gnathostomata) revealed by synchrotron X-ray tomographic microscopy. <i>Palaeoworld</i> , 2019, 28, 441-460.	1.1	11
48	Tubular microfossils from the Ediacaran Wengâ€™an Biota (Doushantuo Formation, South China) are not early animals. <i>Palaeoworld</i> , 2019, 28, 469-477.	1.1	6
49	Anatomy of the Ediacaran rangeomorph <i>Charnia masoni</i> . <i>Papers in Palaeontology</i> , 2019, 5, 157-176.	1.5	16
50	The Early Ediacaran <i>Caveasphaera</i> Foreshadows the Evolutionary Origin of Animal-like Embryology. <i>Current Biology</i> , 2019, 29, 4307-4314.e2.	3.9	16
51	Tooth replacement in early sarcopterygians. <i>Royal Society Open Science</i> , 2019, 6, 191173.	2.4	13
52	Nuclear protein phylogenies support the monophyly of the three bryophyte groups (Bryophyta) Tj ETQq0 0 0 rgBT /Overlock 1.2 Tf 50 1484		
53	The timescale of early land plant evolution. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E2274-E2283.	7.1	654
54	Empirical realism of simulated data is more important than the model used to generate it: a reply to Goloboff & etÂal.. <i>Palaeontology</i> , 2018, 61, 631-635.	2.2	29

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55	Experimental analysis of soft-tissue fossilization: opening the black box. <i>Palaeontology</i> , 2018, 61, 317-323.	2.2	45
56	Constraining uncertainty in the timescale of angiosperm evolution and the veracity of a Cretaceous Terrestrial Revolution. <i>New Phytologist</i> , 2018, 218, 819-834.	7.3	149
57	The Interrelationships of Land Plants and the Nature of the Ancestral Embryophyte. <i>Current Biology</i> , 2018, 28, 733-745.e2.	3.9	398
58	The Efficacy of Consensus Tree Methods for Summarizing Phylogenetic Relationships from a Posterior Sample of Trees Estimated from Morphological Data. <i>Systematic Biology</i> , 2018, 67, 354-362.	5.6	45
59	Probabilistic methods surpass parsimony when assessing clade support in phylogenetic analyses of discrete morphological data. <i>Palaeontology</i> , 2018, 61, 105-118.	2.2	61
60	Ediacaran developmental biology. <i>Biological Reviews</i> , 2018, 93, 914-932.	10.4	80
61	Evolution of jaw disparity in fishes. <i>Palaeontology</i> , 2018, 61, 847-854.	2.2	21
62	Reply to Hedges et al.: Accurate timetrees do indeed require accurate calibrations. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E9512-E9513.	7.1	15
63	Unicellular Origin of the Animal MicroRNA Machinery. <i>Current Biology</i> , 2018, 28, 3288-3295.e5.	3.9	42
64	Evolution of metazoan morphological disparity. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E8909-E8918.	7.1	78
65	Well-Annotated microRNAomes Do Not Evidence Pervasive miRNA Loss. <i>Genome Biology and Evolution</i> , 2018, 10, 1457-1470.	2.5	41
66	The nature of aspidin and the evolutionary origin of bone. <i>Nature Ecology and Evolution</i> , 2018, 2, 1501-1506.	7.8	28
67	Whole-Genome Duplication and Plant Macroevolution. <i>Trends in Plant Science</i> , 2018, 23, 933-945.	8.8	244
68	Integrated genomic and fossil evidence illuminates life's early evolution and eukaryote origin. <i>Nature Ecology and Evolution</i> , 2018, 2, 1556-1562.	7.8	274
69	MicroRNA annotation of plant genomes â˜ Do it right or not at all. <i>BioEssays</i> , 2017, 39, 1600113.	2.5	50
70	Uncertain-tree: discriminating among competing approaches to the phylogenetic analysis of phenotype data. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2017, 284, 20162290.	2.6	114
71	Open data and digital morphology. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2017, 284, 20170194.	2.6	103
72	The Weng'an Biota (Doushantuo Formation): an Ediacaran window on soft-bodied and multicellular microorganisms. <i>Journal of the Geological Society</i> , 2017, 174, 793-802.	2.1	43

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73	The origin of animals: Can molecular clocks and the fossil record be reconciled?. <i>BioEssays</i> , 2017, 39, 1-12.	2.5	105
74	Testing the molecular clock using mechanistic models of fossil preservation and molecular evolution. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2017, 284, 20170227.	2.6	51
75	RelTime Rates Collapse to a Strict Clock When Estimating the Timeline of Animal Diversification. <i>Genome Biology and Evolution</i> , 2017, 9, 1320-1328.	2.5	25
76	Evolution: Divining the Nature of the Ancestral Vertebrate. <i>Current Biology</i> , 2017, 27, R277-R279.	3.9	8
77	Parsimony and maximum-likelihood phylogenetic analyses of morphology do not generally integrate uncertainty in inferring evolutionary history: a response to Brown <i>et al.</i> . <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2017, 284, 20171636.	2.6	19
78	Nuclei and nucleoli in embryo-like fossils from the Ediacaran Weng'an Biota. <i>Precambrian Research</i> , 2017, 301, 145-151.	2.7	30
79	The early Cambrian fossil embryo <i>Pseudoooides</i> is a direct-developing cnidarian, not an early ecdysozoan. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2017, 284, 20172188.	2.6	19
80	Constraining the timing of whole genome duplication in plant evolutionary history. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2017, 284, 20170912.	2.6	47
81	Tips and nodes are complementary not competing approaches to the calibration of molecular clocks. <i>Biology Letters</i> , 2016, 12, 20150975.	2.3	42
82	Bayesian methods outperform parsimony but at the expense of precision in the estimation of phylogeny from discrete morphological data. <i>Biology Letters</i> , 2016, 12, 20160081.	2.3	160
83	Pigmented anatomy in Carboniferous cyclostomes and the evolution of the vertebrate eye. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2016, 283, 20161151.	2.6	44
84	Developmental biology of the early Cambrian cnidarian <i>Olivoooides</i> . <i>Palaeontology</i> , 2016, 59, 387-407.	2.2	29
85	A multicellular organism with embedded cell clusters from the Ediacaran Weng'an biota (Doushantuo Formation, South China). <i>Evolution & Development</i> , 2016, 18, 308-316.	2.0	5
86	Evolution of the Calcium-Based Intracellular Signaling System. <i>Genome Biology and Evolution</i> , 2016, 8, 2118-2132.	2.5	35
87	Reply to 'placoderms and the evolutionary origin of teeth': Burrow <i>et al.</i> (2016). <i>Biology Letters</i> , 2016, 12, 20160526.	2.3	3
88	Dating species divergences using rocks and clocks. <i>Philosophical Transactions of the Royal Society B: Biological Sciences</i> , 2016, 371, 20150126.	4.0	8
89	The evolution of methods for establishing evolutionary timescales. <i>Philosophical Transactions of the Royal Society B: Biological Sciences</i> , 2016, 371, 20160020.	4.0	79
90	Tectonic blocks and molecular clocks. <i>Philosophical Transactions of the Royal Society B: Biological Sciences</i> , 2016, 371, 20160098.	4.0	46

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91	The ins and outs of the evolutionary origin of teeth. <i>Evolution & Development</i> , 2016, 18, 19-30.	2.0	60
92	The Interrelationships of Placental Mammals and the Limits of Phylogenetic Inference. <i>Genome Biology and Evolution</i> , 2016, 8, 330-344.	2.5	195
93	Translating taxonomy into the evolution of conodont feeding ecology. <i>Geology</i> , 2016, 44, 247-250.	4.4	30
94	Histology and affinity of anaspids, and the early evolution of the vertebrate dermal skeleton. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2016, 283, 20152917.	2.6	44
95	Bayesian molecular clock dating of species divergences in the genomics era. <i>Nature Reviews Genetics</i> , 2016, 17, 71-80.	16.3	244
96	Histology of the heterostracan dermal skeleton: Insight into the origin of the vertebrate mineralised skeleton. <i>Journal of Morphology</i> , 2015, 276, 657-680.	1.2	35
97	Cyanobacteria and the Great Oxidation Event: evidence from genes and fossils. <i>Palaeontology</i> , 2015, 58, 769-785.	2.2	207
98	Experimental taphonomy of <i>Artemia</i> reveals the role of endogenous microbes in mediating decay and fossilization. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2015, 282, 20150476.	2.6	65
99	Cyanobacteria and the Great Oxidation Event: evidence from genes and fossils. <i>Palaeontology</i> , 2015, 58, 935-936.	2.2	8
100	Size is not everything: rates of genome size evolution, not <i>C</i> -value, correlate with speciation in angiosperms. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2015, 282, 20152289.	2.6	65
101	Calibration uncertainty in molecular dating analyses: there is no substitute for the prior evaluation of time priors. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2015, 282, 20141013.	2.6	184
102	<i>Romundina</i> and the evolutionary origin of teeth. <i>Biology Letters</i> , 2015, 11, 20150326.	2.3	28
103	Critical appraisal of tubular putative eumetazoans from the Ediacaran Weng'an Doushantuo biota. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2015, 282, 20151169.	2.6	21
104	Constraining the Deep Origin of Parasitic Flatworms and Host-Interactions with Fossil Evidence. <i>Advances in Parasitology</i> , 2015, 90, 93-135.	3.2	47
105	The Fossil Calibration Database—A New Resource for Divergence Dating. <i>Systematic Biology</i> , 2015, 64, 853-859.	5.6	54
106	Do cladistic and morphometric data capture common patterns of morphological disparity?. <i>Palaeontology</i> , 2015, 58, 393-399.	2.2	45
107	Ptychographic nanotomography at the Swiss Light Source. <i>Proceedings of SPIE</i> , 2015, , .	0.8	2
108	Uncertainty in the Timing of Origin of Animals and the Limits of Precision in Molecular Timescales. <i>Current Biology</i> , 2015, 25, 2939-2950.	3.9	370

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109	Dating Tips for Divergence-Time Estimation. <i>Trends in Genetics</i> , 2015, 31, 637-650.	6.7	126
110	Embryology in Deep Time. , 2015, , 45-63.		11
111	Discriminating signal from noise in the fossil record of early vertebrates reveals cryptic evolutionary history. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2015, 282, 20142245.	2.6	31
112	Distinguishing Biology from Geology in Soft-Tissue Preservation. <i>The Paleontological Society Papers</i> , 2014, 20, 275-288.	0.6	7
113	Evaluating scenarios for the evolutionary assembly of the brachiopod body plan. <i>Evolution & Development</i> , 2014, 16, 13-24.	2.0	22
114	Functional adaptation underpinned the evolutionary assembly of the earliest vertebrate skeleton. <i>Evolution & Development</i> , 2014, 16, 354-361.	2.0	9
115	Evolutionary history of plant microRNAs. <i>Trends in Plant Science</i> , 2014, 19, 175-182.	8.8	182
116	Finite element, occlusal, microwear and microstructural analyses indicate that conodont microstructure is adapted to dental function. <i>Palaeontology</i> , 2014, 57, 1059-1066.	2.2	30
117	Neither phylogenomic nor palaeontological data support a Palaeogene origin of placental mammals. <i>Biology Letters</i> , 2014, 10, 20131003.	2.3	87
118	Early vertebrate evolution. <i>Palaeontology</i> , 2014, 57, 879-893.	2.2	56
119	Developmental paleobiology of the vertebrate skeleton. <i>Journal of Paleontology</i> , 2014, 88, 676-683.	0.8	12
120	There is no general model for occlusal kinematics in conodonts. <i>Lethaia</i> , 2014, 47, 547-555.	1.4	16
121	A virtual world of paleontology. <i>Trends in Ecology and Evolution</i> , 2014, 29, 347-357.	8.7	205
122	A divergence dating analysis of turtles using fossil calibrations: an example of best practices. <i>Journal of Paleontology</i> , 2013, 87, 612-634.	0.8	128
123	Histology of <i>œplacodermâ•</i> dermal skeletons: Implications for the nature of the ancestral gnathostome. <i>Journal of Morphology</i> , 2013, 274, 627-644.	1.2	58
124	Embryos, polyps and medusae of the Early Cambrian scyphozoan <i>Olivoooides</i> . <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2013, 280, 20130071.	2.6	66
125	A daily-updated tree of (sequenced) life as a reference for genome research. <i>Scientific Reports</i> , 2013, 3, 2015.	3.3	47
126	Cutting the first â€˜teethâ€™: a new approach to functional analysis of conodont elements. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2013, 280, 20131524.	2.6	13

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127	The origin of conodonts and of vertebrate mineralized skeletons. <i>Nature</i> , 2013, 502, 546-549.	27.8	79
128	miRNAs: Small Genes with Big Potential in Metazoan Phylogenetics. <i>Molecular Biology and Evolution</i> , 2013, 30, 2369-2382.	8.9	118
129	Response to Comment on ‘Fossilized Nuclei and Germination Structures Identify Ediacaran ‘Animal Embryos’ as Encysting Protists’. <i>Science</i> , 2012, 335, 1169-1169.	12.6	14
130	Testing models of dental development in the earliest bony vertebrates, <i>Andreolepis</i> and <i>Lophosteus</i> . <i>Biology Letters</i> , 2012, 8, 833-837.	2.3	18
131	Experimental taphonomy of giant sulphur bacteria: implications for the interpretation of the embryo-like Ediacaran Doushantuo fossils. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2012, 279, 1857-1864.	2.6	45
132	Phylogenomic datasets provide both precision and accuracy in estimating the timescale of placental mammal phylogeny. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2012, 279, 3491-3500.	2.6	449
133	A merciful death for the ‘earliest bilaterian’, <i>Vernanimalcula</i> . <i>Evolution & Development</i> , 2012, 14, 421-427.	2.0	33
134	Development of teeth and jaws in the earliest jawed vertebrates. <i>Nature</i> , 2012, 491, 748-751.	27.8	98
135	Do miRNAs have a deep evolutionary history?. <i>BioEssays</i> , 2012, 34, 857-866.	2.5	96
136	Ontogeny and microstructure of the enigmatic Cambrian tommotiid <i>< i>Sunnaginia</i></i> Missarzhevsky, 1969. <i>Palaeontology</i> , 2012, 55, 661-676.	2.2	26
137	New palaeoscolecid worms from the Furongian (upper Cambrian) of Hunan, South China: is <i>< i>Markuelia</i></i> an embryonic palaeoscolecid?. <i>Palaeontology</i> , 2012, 55, 613-622.	2.2	25
138	Exploring uncertainty in the calibration of the molecular clock. <i>Biology Letters</i> , 2012, 8, 156-159.	2.3	206
139	Testing microstructural adaptation in the earliest dental tools. <i>Biology Letters</i> , 2012, 8, 952-955.	2.3	15
140	Distinguishing geology from biology in the Ediacaran Doushantuo biota relaxes constraints on the timing of the origin of bilaterians. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2012, 279, 2369-2376.	2.6	43
141	The sharpest tools in the box? Quantitative analysis of conodont element functional morphology. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2012, 279, 2849-2854.	2.6	49
142	Best Practices for Justifying Fossil Calibrations. <i>Systematic Biology</i> , 2012, 61, 346-359.	5.6	616
143	Evolutionary crossroads in developmental biology: cyclostomes (lamprey and hagfish). <i>Development (Cambridge)</i> , 2012, 139, 2091-2099.	2.5	142
144	Fossilized Nuclei and Germination Structures Identify Ediacaran ‘Animal Embryos’ as Encysting Protists. <i>Science</i> , 2011, 334, 1696-1699.	12.6	142

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145	Fossil jawless fish from China foreshadows early jawed vertebrate anatomy. <i>Nature</i> , 2011, 476, 324-327.	27.8	112
146	Evolutionary Origins of Animal Skeletal Biominerization. <i>Cells Tissues Organs</i> , 2011, 194, 98-102.	2.3	74
147	Response by Philip Donoghue. <i>Journal of Paleontology</i> , 2011, 85, 1016-1016.	0.8	0
148	Teeth before jaws? Comparative analysis of the structure and development of the external and internal scales in the extinct jawless vertebrate <i>Laganellia scotica</i> . <i>Evolution & Development</i> , 2011, 13, 523-532.	2.0	34
149	Establishing a time-scale for plant evolution. <i>New Phytologist</i> , 2011, 192, 266-301.	7.3	306
150	Is evolutionary history repeatedly rewritten in light of new fossil discoveries?. <i>Proceedings of the Royal Society B: Biological Sciences</i> , 2011, 278, 599-604.	2.6	16
151	The Trouble with Topology: Phylogenies without Fossils Provide a Revisionist Perspective of Evolutionary History in Topological Analyses of Diversity. <i>Systematic Biology</i> , 2011, 60, 700-712.	5.6	20
152	Are palaeoscolecids ancestral ecdysozoans?. <i>Evolution & Development</i> , 2010, 12, 177-200.	2.0	83
153	The anatomy, taphonomy, taxonomy and systematic affinity of Markuelia: Early Cambrian to Early Ordovician scalidophorans. <i>Palaeontology</i> , 2010, 53, 1291-1314.	2.2	53
154	Origins of multicellularity. <i>Nature</i> , 2010, 466, 41-42.	27.8	51
155	A formula for maximum possible steps in multistate characters: isolating matrix parameter effects on measures of evolutionary convergence. <i>Cladistics</i> , 2010, 26, 98-102.	3.3	11
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