

# J H Duncan Bassett

## List of Publications by Year in Descending Order

**Source:** <https://exaly.com/author-pdf/7572089/j-h-duncan-bassett-publications-by-year.pdf>

**Version:** 2024-04-19

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

79  
papers

3,977  
citations

36  
h-index

62  
g-index

85  
ext. papers

5,039  
ext. citations

7.8  
avg, IF

5.65  
L-index

#	Paper	IF	Citations
79	Bone Phenotyping Approaches in Human, Mice and Zebrafish - Expert Overview of the EU Cost Action GEMSTONE ("GENomics of MusculoSkeletal traits TranslatiOnal NETwork").. <i>Frontiers in Endocrinology</i> , <b>2021</b> , 12, 720728	5.7	0
78	Osteoclasts recycle via osteomorphs during RANKL-stimulated bone resorption. <i>Cell</i> , <b>2021</b> , 184, 1330-1347	17.4	139
77	Osteocyte transcriptome mapping identifies a molecular landscape controlling skeletal homeostasis and susceptibility to skeletal disease. <i>Nature Communications</i> , <b>2021</b> , 12, 2444	17.4	12
76	An variant links aberrant Rac1 function to early-onset skeletal fragility. <i>JBMR Plus</i> , <b>2021</b> , 5, e10509	3.9	
75	Thyroid hormone, thyroid medication, and the skeleton <b>2021</b> , 1139-1157		
74	A molecular quantitative trait locus map for osteoarthritis. <i>Nature Communications</i> , <b>2021</b> , 12, 1309	17.4	8
73	A Roadmap to Gene Discoveries and Novel Therapies in Monogenic Low and High Bone Mass Disorders. <i>Frontiers in Endocrinology</i> , <b>2021</b> , 12, 709711	5.7	3
72	Accelerating functional gene discovery in osteoarthritis. <i>Nature Communications</i> , <b>2021</b> , 12, 467	17.4	12
71	A Polygenic Risk Score as a Risk Factor for Medication-Associated Fractures. <i>Journal of Bone and Mineral Research</i> , <b>2020</b> , 35, 1935-1941	6.3	3
70	Response to Letter to the Editor: "IGSF1 Deficiency Results in Human and Murine Somatotrope Neurosecretory Hyperfunction". <i>Journal of Clinical Endocrinology and Metabolism</i> , <b>2020</b> , 105,	5.6	
69	Role of thyroid hormones in craniofacial development. <i>Nature Reviews Endocrinology</i> , <b>2020</b> , 16, 147-164	15.2	14
68	Mouse mutant phenotyping at scale reveals novel genes controlling bone mineral density. <i>PLoS Genetics</i> , <b>2020</b> , 16, e1009190	6	8
67	A trans-eQTL network regulates osteoclast multinucleation and bone mass. <i>ELife</i> , <b>2020</b> , 9,	8.9	10
66	IGSF1 Deficiency Results in Human and Murine Somatotrope Neurosecretory Hyperfunction. <i>Journal of Clinical Endocrinology and Metabolism</i> , <b>2020</b> , 105,	5.6	14
65	Quantitative X-Ray Imaging of Mouse Bone by Faxitron. <i>Methods in Molecular Biology</i> , <b>2019</b> , 1914, 559-569	5.9	5
64	Slc20a2, Encoding the Phosphate Transporter PiT2, Is an Important Genetic Determinant of Bone Quality and Strength. <i>Journal of Bone and Mineral Research</i> , <b>2019</b> , 34, 1101-1114	6.3	18
63	PYY is a negative regulator of bone mass and strength. <i>Bone</i> , <b>2019</b> , 127, 427-435	4.7	6

62	Type 2 deiodinase polymorphism causes ER stress and hypothyroidism in the brain. <i>Journal of Clinical Investigation</i> , <b>2019</b> , 129, 230-245	15.9	44
61	An atlas of genetic influences on osteoporosis in humans and mice. <i>Nature Genetics</i> , <b>2019</b> , 51, 258-266	36.3	270
60	Transferrin receptor 2 controls bone mass and pathological bone formation via BMP and Wnt signaling. <i>Nature Metabolism</i> , <b>2019</b> , 1, 111-124	14.6	36
59	The bone remodelling cycle. <i>Annals of Clinical Biochemistry</i> , <b>2018</b> , 55, 308-327	2.2	179
58	Life-Course Genome-wide Association Study Meta-analysis of Total Body BMD and Assessment of Age-Specific Effects. <i>American Journal of Human Genetics</i> , <b>2018</b> , 102, 88-102	11	119
57	Frequent falls and confusion: recurrent hypoglycemia in a patient with tuberous sclerosis complex. <i>Clinical Case Reports (discontinued)</i> , <b>2018</b> , 6, 904-909	0.7	3
56	Thyroid Stimulating Hormone and Bone Mineral Density: Evidence From a Two-Sample Mendelian Randomization Study and a Candidate Gene Association Study. <i>Journal of Bone and Mineral Research</i> , <b>2018</b> , 33, 1318-1325	6.3	18
55	Thyroid diseases and bone health. <i>Journal of Endocrinological Investigation</i> , <b>2018</b> , 41, 99-109	5.2	78
54	Common signalling pathways in macrophage and osteoclast multinucleation. <i>Journal of Cell Science</i> , <b>2018</b> , 131,	5.3	83
53	Genome-wide association study of extreme high bone mass: Contribution of common genetic variation to extreme BMD phenotypes and potential novel BMD-associated genes. <i>Bone</i> , <b>2018</b> , 114, 62-74	4.7	25
52	Analysis of Physiological Responses to Thyroid Hormones and Their Receptors in Bone. <i>Methods in Molecular Biology</i> , <b>2018</b> , 1801, 123-154	1.4	8
51	Animal Models <b>2018</b> , 359-366		
50	Thyroid Hormone in Bone and Joint Disorders <b>2018</b> , 547-569		
49	Inhibiting the osteocyte-specific protein sclerostin increases bone mass and fracture resistance in multiple myeloma. <i>Blood</i> , <b>2017</b> , 129, 3452-3464	2.2	117
48	An Essential Physiological Role for MCT8 in Bone in Male Mice. <i>Endocrinology</i> , <b>2017</b> , 158, 3055-3066	4.8	11
47	Noncanonical thyroid hormone signaling mediates cardiometabolic effects in vivo. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2017</b> , 114, E11323-E11332	11.5	53
46	Identification of 153 new loci associated with heel bone mineral density and functional involvement of GPC6 in osteoporosis. <i>Nature Genetics</i> , <b>2017</b> , 49, 1468-1475	36.3	235
45	Role of Thyroid Hormones in Skeletal Development and Bone Maintenance. <i>Endocrine Reviews</i> , <b>2016</b> , 37, 135-87	27.2	217

44	Rapid phenotyping of knockout mice to identify genetic determinants of bone strength. <i>Journal of Endocrinology</i> , <b>2016</b> , 231, R31-46	4.7	24
43	An undiagnosed stupor in the acute medical unit: a case of malignant catatonia. <i>QJM - Monthly Journal of the Association of Physicians</i> , <b>2015</b> , 108, 335-6	2.7	1
42	Adult mice lacking the type 2 iodothyronine deiodinase have increased subchondral bone but normal articular cartilage. <i>Thyroid</i> , <b>2015</b> , 25, 269-77	6.2	14
41	Thyrostimulin Regulates Osteoblastic Bone Formation During Early Skeletal Development. <i>Endocrinology</i> , <b>2015</b> , 156, 3098-113	4.8	33
40	Classification and proposed nomenclature for inherited defects of thyroid hormone action, cell transport, and metabolism. <i>Journal of Clinical Endocrinology and Metabolism</i> , <b>2014</b> , 99, 768-70	5.6	44
39	Kcnn4 is a regulator of macrophage multinucleation in bone homeostasis and inflammatory disease. <i>Cell Reports</i> , <b>2014</b> , 8, 1210-24	10.6	41
38	Classification and proposed nomenclature for inherited defects of thyroid hormone action, cell transport, and metabolism. <i>Thyroid</i> , <b>2014</b> , 24, 407-9	6.2	37
37	Quantitative X-ray microradiography for high-throughput phenotyping of osteoarthritis in mice. <i>Osteoarthritis and Cartilage</i> , <b>2014</b> , 22, 1396-400	6.2	12
36	Thyroid hormone receptor $\beta$ mutation causes a severe and thyroxine-resistant skeletal dysplasia in female mice. <i>Endocrinology</i> , <b>2014</b> , 155, 3699-712	4.8	41
35	Mechanisms of action of thyroid hormones in the skeleton. <i>Biochimica Et Biophysica Acta - General Subjects</i> , <b>2013</b> , 1830, 3979-86	4	64
34	Thyroid hormone metabolism in skeletal development and adult bone maintenance. <i>Trends in Endocrinology and Metabolism</i> , <b>2012</b> , 23, 155-62	8.8	67
33	Significant deterioration in nanomechanical quality occurs through incomplete extrafibrillar mineralization in rachitic bone: evidence from in-situ synchrotron X-ray scattering and backscattered electron imaging. <i>Journal of Bone and Mineral Research</i> , <b>2012</b> , 27, 876-90	6.3	53
32	Genetic evidence that thyroid hormone is indispensable for prepubertal insulin-like growth factor-I expression and bone acquisition in mice. <i>Journal of Bone and Mineral Research</i> , <b>2012</b> , 27, 1067-79	6.3	62
31	A mouse model for spondyloepiphyseal dysplasia congenita with secondary osteoarthritis due to a Col2a1 mutation. <i>Journal of Bone and Mineral Research</i> , <b>2012</b> , 27, 413-28	6.3	27
30	Rapid-throughput skeletal phenotyping of 100 knockout mice identifies 9 new genes that determine bone strength. <i>PLoS Genetics</i> , <b>2012</b> , 8, e1002858	6	62
29	The skeletal consequences of thyrotoxicosis. <i>Journal of Endocrinology</i> , <b>2012</b> , 213, 209-21	4.7	73
28	Mice lacking the calcineurin inhibitor Rcan2 have an isolated defect of osteoblast function. <i>Endocrinology</i> , <b>2012</b> , 153, 3537-48	4.8	20
27	Quantitative X-ray imaging of rodent bone by Faxitron. <i>Methods in Molecular Biology</i> , <b>2012</b> , 816, 499-506	4.4	25

26	Bone Mineral Content and Density. <i>Current Protocols in Mouse Biology</i> , <b>2012</b> , 2, 365-400	1.1	5
25	Deiodinases: the balance of thyroid hormone: local control of thyroid hormone action: role of type 2 deiodinase. <i>Journal of Endocrinology</i> , <b>2011</b> , 209, 261-72	4.7	89
24	Optimal bone strength and mineralization requires the type 2 iodothyronine deiodinase in osteoblasts. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2010</b> , 107, 7604-9	11.5	106
23	Thyroid and bone. <i>Archives of Biochemistry and Biophysics</i> , <b>2010</b> , 503, 129-36	4.1	112
22	The skeletal phenotypes of TRalpha and TRbeta mutant mice. <i>Journal of Molecular Endocrinology</i> , <b>2009</b> , 42, 269-82	4.5	63
21	Bone signaling pathways and treatment of osteoporosis. <i>Expert Review of Endocrinology and Metabolism</i> , <b>2009</b> , 4, 639-650	4.1	11
20	Critical role of the hypothalamic-pituitary-thyroid axis in bone. <i>Bone</i> , <b>2008</b> , 43, 418-26	4.7	94
19	A lack of thyroid hormones rather than excess thyrotropin causes abnormal skeletal development in hypothyroidism. <i>Molecular Endocrinology</i> , <b>2008</b> , 22, 501-12		94
18	Thyroid status during skeletal development determines adult bone structure and mineralization. <i>Molecular Endocrinology</i> , <b>2007</b> , 21, 1893-904		102
17	Thyroid hormone excess rather than thyrotropin deficiency induces osteoporosis in hyperthyroidism. <i>Molecular Endocrinology</i> , <b>2007</b> , 21, 1095-107		123
16	Thyroid hormone regulates heparan sulfate proteoglycan expression in the growth plate. <i>Endocrinology</i> , <b>2006</b> , 147, 295-305	4.8	36
15	Characterization of skeletal phenotypes of TRalpha1 and TRbeta mutant mice: implications for tissue thyroid status and T3 target gene expression. <i>Nuclear Receptor Signaling</i> , <b>2006</b> , 4, e011	1	38
14	Analysis of skeletal phenotypes in thyroid hormone receptor mutant mice. <i>Scanning</i> , <b>2006</b> , 28, 91-93	1.6	6
13	Thyroid hormones regulate fibroblast growth factor receptor signaling during chondrogenesis. <i>Endocrinology</i> , <b>2005</b> , 146, 5568-80	4.8	66
12	Contrasting skeletal phenotypes in mice with an identical mutation targeted to thyroid hormone receptor alpha1 or beta. <i>Molecular Endocrinology</i> , <b>2005</b> , 19, 3045-59		109
11	A Tense Case—Turney’s Triad. <i>Journal of the Royal Society of Medicine</i> , <b>2004</b> , 97, 540-541	2.3	
10	Mechanisms of thyroid hormone receptor-specific nuclear and extra nuclear actions. <i>Molecular and Cellular Endocrinology</i> , <b>2003</b> , 213, 1-11	4.4	288
9	The molecular actions of thyroid hormone in bone. <i>Trends in Endocrinology and Metabolism</i> , <b>2003</b> , 14, 356-64	8.8	184

8	Novel DAX1 mutations in X-linked adrenal hypoplasia congenita and hypogonadotropic hypogonadism. <i>Clinical Endocrinology</i> , <b>1999</b> , 50, 69-75	3.4	30
7	Studies of the murine homolog of the multiple endocrine neoplasia type 1 (MEN1) gene, men1. <i>Journal of Bone and Mineral Research</i> , <b>1999</b> , 14, 3-10	6.3	39
6	Mapping of the gene encoding the B56 beta subunit of protein phosphatase 2A (PPP2R5B) to a 0.5-Mb region of chromosome 11q13 and its exclusion as a candidate gene for multiple endocrine neoplasia type 1 (MEN1). <i>Human Genetics</i> , <b>1997</b> , 100, 481-5	6.3	1
5	The European Consortium on MEN1. Linkage disequilibrium studies in multiple endocrine neoplasia type 1 (MEN1). <i>Human Genetics</i> , <b>1997</b> , 100, 657-65	6.3	9
4	Accelerating functional gene discovery in osteoarthritis		1
3	Osteocyte Transcriptome Mapping Identifies a Molecular Landscape Controlling Skeletal Homeostasis and Susceptibility to Skeletal Disease		3
2	An Atlas of Human and Murine Genetic Influences on Osteoporosis		3
1	Decoding the genomic basis of osteoarthritis		4