

# Igor Fijalkowski

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3933811/publications.pdf>

Version: 2024-02-01

16  
papers

301  
citations

1040056

9  
h-index

1125743

13  
g-index

17  
all docs

17  
docs citations

17  
times ranked

517  
citing authors

#	ARTICLE	IF	CITATIONS
1	A Novel Domain-Specific Mutation in a Sclerosteosis Patient Suggests a Role of LRP4 as an Anchor for Sclerostin in Human Bone. <i>Journal of Bone and Mineral Research</i> , 2016, 31, 874-881.	2.8	65
2	The role of extracellular modulators of canonical Wnt signaling in bone metabolism and diseases. <i>Seminars in Arthritis and Rheumatism</i> , 2013, 43, 220-240.	3.4	62
3	Genetic control of bone mass. <i>Molecular and Cellular Endocrinology</i> , 2016, 432, 3-13.	3.2	59
4	The <i>Lrp4</i> R1170Q Homozygous Knock-In Mouse Recapitulates the Bone Phenotype of Sclerosteosis in Humans. <i>Journal of Bone and Mineral Research</i> , 2017, 32, 1739-1749.	2.8	27
5	Lost and Found: Re-searching and Re-scoring Proteomics Data Aids Genome Annotation and Improves Proteome Coverage. <i>MSystems</i> , 2020, 5, .	3.8	13
6	Sclerosing Bone Dysplasias: Leads Toward Novel Osteoporosis Treatments. <i>Current Osteoporosis Reports</i> , 2014, 12, 243-251.	3.6	12
7	Bacterial riboproteogenomics: the era of N-terminal proteoform existence revealed. <i>FEMS Microbiology Reviews</i> , 2020, 44, 418-431.	8.6	12
8	Small Protein Enrichment Improves Proteomics Detection of sORF Encoded Polypeptides. <i>Frontiers in Genetics</i> , 2021, 12, 713400.	2.3	12
9	Hidden in plain sight: challenges in proteomics detection of small ORF-encoded polypeptides. <i>MicroLife</i> , 2022, 3, .	2.1	10
10	Eight mutations including 5 novel ones in the COL1A1 gene in Czech patients with osteogenesis imperfecta. <i>Biomedical Papers of the Medical Faculty of the University Palacky, Olomouc, Czechoslovakia</i> , 2016, 160, 442-447.	0.6	9
11	Further delineation of facioaudiosymphalangism syndrome: Description of a family with a novel <i>NOC</i> mutation and without hearing loss. <i>American Journal of Medical Genetics, Part A</i> , 2016, 170, 1479-1484.	1.2	8
12	Mutations in sFRP1 or sFRP4 are not a common cause of craniotubular hyperostosis. <i>Bone</i> , 2013, 52, 292-295.	2.9	4
13	Capturing Salmonella SspH2 Host Targets in Virus-Like Particles. <i>Frontiers in Medicine</i> , 2021, 8, 725072.	2.6	4
14	Expression analysis of mesenchymal KS483 cells during differentiation towards osteoblasts. <i>Bone Abstracts</i> , 0, , .	0.0	0
15	Rs55710688 in the Kozak sequence of WNT16 increases translation efficiency and is associated with osteoporosis related parameters. <i>Bone Abstracts</i> , 0, , .	0.0	0
16	Common variants in <i>Rspo 1,2 and 3</i> do not associate with BMD in stratified subpopulations of the Odense Androgen Study and mutations in these genes are not a common cause of craniotubular hyperostosis. <i>Bone Abstracts</i> , 0, , .	0.0	0