

# Nerea Alonso

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6429157/publications.pdf>

Version: 2024-02-01

25  
papers

2,478  
citations

516561

16  
h-index

610775

24  
g-index

25  
all docs

25  
docs citations

25  
times ranked

4914  
citing authors

#	ARTICLE	IF	CITATIONS
1	Pattern of SQSTM1 Gene Variants in a Hungarian Cohort of Paget's Disease of Bone. <i>Calcified Tissue International</i> , 2021, 108, 159-164.	1.5	2
2	Review of Current Real-World Experience with Teriparatide as Treatment of Osteoporosis in Different Patient Groups. <i>Journal of Clinical Medicine</i> , 2021, 10, 1403.	1.0	12
3	The "Genomics of Musculo Skeletal Traits Translational Network": Origins, Rationale, Organization, and Prospects. <i>Frontiers in Endocrinology</i> , 2021, 12, 709815.	1.5	3
4	Loss-of-Function Mutations in the <i>ALPL</i> Gene Presenting with Adult Onset Osteoporosis and Low Serum Concentrations of Total Alkaline Phosphatase. <i>Journal of Bone and Mineral Research</i> , 2020, 35, 657-661.	3.1	23
5	Adult hypophosphatasia with a novel ALPL mutation: Report of an Indian kindred. <i>Bone Reports</i> , 2020, 12, 100247.	0.2	3
6	Opportunities and Challenges in Functional Genomics Research in Osteoporosis: Report From a Workshop Held by the Causes Working Group of the Osteoporosis and Bone Research Academy of the Royal Osteoporosis Society on October 5th 2020. <i>Frontiers in Endocrinology</i> , 2020, 11, 630875.	1.5	5
7	Insertion Mutation in <i>Tnfrsf11a</i> Causes a Paget's Disease-Like Phenotype in Heterozygous Mice and Osteopetrosis in Homozygous Mice. <i>Journal of Bone and Mineral Research</i> , 2020, 36, 1376-1386.	3.1	10
8	Genetics of Osteoporosis. , 2020, , 83-91.		0
9	Novel clinical and molecular findings in Spanish patients with naevoid basal cell carcinoma syndrome. <i>British Journal of Dermatology</i> , 2018, 178, 198-206.	1.4	5
10	Identification of a novel locus on chromosome 2q13, which predisposes to clinical vertebral fractures independently of bone density. <i>Annals of the Rheumatic Diseases</i> , 2018, 77, 378-385.	0.5	21
11	Clinical and Genetic Advances in Paget's Disease of Bone: a Review. <i>Clinical Reviews in Bone and Mineral Metabolism</i> , 2017, 15, 37-48.	1.3	37
12	Novel Genetic Variants Associated With Increased Vertebral Volumetric BMD, Reduced Vertebral Fracture Risk, and Increased Expression of <i>SLC1A3</i> and <i>EPHB2</i> . <i>Journal of Bone and Mineral Research</i> , 2016, 31, 2085-2097.	3.1	42
13	Atypical Femoral Fracture in Osteoporosis Pseudoglioma Syndrome Associated with Two Novel Compound Heterozygous Mutations in <i>LRP5</i> . <i>Journal of Bone and Mineral Research</i> , 2015, 30, 615-620.	3.1	25
14	Big data challenges in bone research: genome-wide association studies and next-generation sequencing. <i>BoneKEY Reports</i> , 2015, 4, 635.	2.7	12
15	Whole-genome sequencing identifies EN1 as a determinant of bone density and fracture. <i>Nature</i> , 2015, 526, 112-117.	13.7	483
16	A genome-wide copy number association study of osteoporotic fractures points to the 6p25.1 locus. <i>Journal of Medical Genetics</i> , 2014, 51, 122-131.	1.5	36
17	Genome-wide association study for radiographic vertebral fractures: A potential role for the 16q24 BMD locus. <i>Bone</i> , 2014, 59, 20-27.	1.4	32
18	Unveiling the mysteries of the genetics of osteoporosis. <i>Journal of Endocrinological Investigation</i> , 2014, 37, 925-934.	1.8	20

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19	Genome-wide association study for radiographic vertebral fractures: a potential role for the 16q24 BMD locus. <i>Bone</i> , 2014, 59, 20-7.	1.4	17
20	Common susceptibility alleles and <i>SQSTM1</i> mutations predict disease extent and severity in a multinational study of patients with Paget's disease. <i>Journal of Bone and Mineral Research</i> , 2013, 28, 2338-2346.	3.1	50
21	Genome-wide meta-analysis identifies 56 bone mineral density loci and reveals 14 loci associated with risk of fracture. <i>Nature Genetics</i> , 2012, 44, 491-501.	9.4	1,100
22	Assessment of gene-by-sex interaction effect on bone mineral density. <i>Journal of Bone and Mineral Research</i> , 2012, 27, 2051-2064.	3.1	47
23	Genome-wide association identifies three new susceptibility loci for Paget's disease of bone. <i>Nature Genetics</i> , 2011, 43, 685-689.	9.4	158
24	Mutations of <i>SQSTM1</i> are associated with severity and clinical outcome in paget disease of bone. <i>Journal of Bone and Mineral Research</i> , 2010, 25, 2368-2373.	3.1	77
25	Genome-wide association study identifies variants at <i>CSF1</i> , <i>OPTN</i> and <i>TNFRSF11A</i> as genetic risk factors for Paget's disease of bone. <i>Nature Genetics</i> , 2010, 42, 520-524.	9.4	258