Weibo Xia

List of Publications by Year
in descending order

133 papers	2,799 citations	236833 25 h-index	233338 45 g-index
145 all docs	145 docs citations	145 times ranked	3399 citing authors

#	Article	IF	Citations
1	A novel long-range deletion spanning CDC73 and upper-stream genes discovered in a kindred of familial primary hyperparathyroidism. Endocrine, 2022, 75, 907-915.	1.1	2
2	A robust method for simultaneous measurement of serum 25(OH)D, 1,25(OH)₂D, and 24,25(OH)₂D by liquid chromatographyâ€tandem mass spectrometry with efficient separation of 3â€epi analogs, 23R,25(OH)₂D₃, and 4Î²,25(OH)₂D₃. Journal of Mass Spectrometry, 2022, 57, e4792.	0.7	3
3	Clinical, Biochemical, Radiological, and Genetic Analyses of a Patient with VCP Gene Variant-Induced Pagetâ€™s Disease of Bone. Calcified Tissue International, 2022, 110, 518-528.	1.5	2
4	The efficacy and safety of different doses of calcitriol combined with neutral phosphate in X-linked hypophosphatemia: a prospective study. Osteoporosis International, 2022, 33, 1385-1395.	1.3	3
5	Bone microarchitecture impairment in prolactinoma patients assessed by HR-pQCT. Osteoporosis International, 2022, 33, 1535-1544.	1.3	2
6	Serum Metabolomics Reveals Dysregulation and Diagnostic Potential of Oxylipins in Tumor-induced Osteomalacia. Journal of Clinical Endocrinology and Metabolism, 2022, 107, 1383-1391.	1.8	5
7	Bone microstructure evaluated by TBS and HR-pQCT in Chinese adults with X-linked hypophosphatemia. Bone, 2022, 160, 116423.	1.4	4
8	Overview of the clinical efficacy and safety of eldecalcitol for the treatment of osteoporosis. Archives of Osteoporosis, 2022, 17, 74.	1.0	2
9	Relationship of Pathogenic Mutations and Responses to Zoledronic Acid in a Cohort of Osteogenesis Imperfecta Children. Journal of Clinical Endocrinology and Metabolism, 2022, 107, 2571-2579.	1.8	8
10	Clinical Characteristics and Surgical Outcomes of Sinonasal Lesions Associated With Tumorâ€Induced Osteomalacia. Otolaryngology - Head and Neck Surgery, 2021, 165, 223-231.	1.1	4
11	Consensus Statement on the Use of Bone Turnover Markers for Short-Term Monitoring of Osteoporosis Treatment in the Asia-Pacific Region. Journal of Clinical Densitometry, 2021, 24, 3-13.	0.5	35
12	Iron deficiency plays essential roles in the trigger, treatment, and prognosis of autosomal dominant hypophosphatemic rickets. Osteoporosis International, 2021, 32, 737-745.	1.3	11
13	Comparison of differences in bone microarchitecture in adult- versus juvenile-onset type 1 diabetes Asian males versus non-diabetes males: an observational cross-sectional pilot study. Endocrine, 2021, 71, 87-95.	1.1	10
14	Locoregional recurrence of parathyroid carcinoma: how to identify this rare but fatal condition using ultrasonography. Endocrine Journal, 2021, 68, 1179-1186.	0.7	1
15	Comparison of Differences in Bone Mineral Density Measurement With 3 Hologic Dual-Energy X-Ray Absorptiometry Scan Modes. Journal of Clinical Densitometry, 2021, 24, 645-650.	0.5	2
16	Cushingâ€™s Syndrome With Nocardiosis: A Case Report and a Systematic Review of the Literature. Frontiers in Endocrinology, 2021, 12, 640998.	1.5	2
17	Associations between Osteocalcin, Calciotropic Hormones, and Energy Metabolism in a Cohort of Chinese Postmenopausal Women: Peking Vertebral Fracture Study. International Journal of Endocrinology, 2021, 2021, 1-10.	0.6	3
18	Bone mineral density and microarchitecture among Chinese patients with rheumatoid arthritis: a cross-sectional study with HRpQCT. Arthritis Research and Therapy, 2021, 23, 127.	1.6	13

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19	Case Report: Two Novel Frameshift Mutations in SLC20A2 and One Novel Splice Donor Mutation in PDGFB Associated With Primary Familial Brain Calcification. Frontiers in Genetics, 2021, 12, 643452.	1.1	4
20	Bone turnover markers are associated with the PTH levels in different subtypes of pseudohypoparathyroidism type 1 patients. Clinical Endocrinology, 2021, 95, 277-285.	1.2	1
21	Exome sequencing reveals genetic architecture in patients with isolated or syndromic short stature. Journal of Genetics and Genomics, 2021, 48, 396-402.	1.7	21
22	Health-related quality of life in men with osteoporosis: a systematic review and meta-analysis. Endocrine, 2021, 74, 270-280.	1.1	12
23	Novel AQP2 Mutations and Clinical Characteristics in Seven Chinese Families With Congenital Nephrogenic Diabetes Insipidus. Frontiers in Endocrinology, 2021, 12, 686818.	1.5	2
24	Comparative effect of eldecalcitol and alfacalcidol on bone microstructure: A preliminary report of secondary analysis of a prospective trial. Osteoporosis and Sarcopenia, 2021, 7, 47-53.	0.7	1
25	Novel AVPR2 mutations and clinical characteristics in 28 Chinese families with congenital nephrogenic diabetes insipidus. Journal of Endocrinological Investigation, 2021, 44, 2777-2783.	1.8	5
26	Low Levels of Serum Sclerostin in Adult Patients With Tumor-Induced Osteomalacia Compared With X-linked Hypophosphatemia. Journal of Clinical Endocrinology and Metabolism, 2021, , .	1.8	2
27	Molecular Characterization of an Aquaporinâ˜2 Mutation Causing Nephrogenic Diabetes Insipidus. Frontiers in Endocrinology, 2021, 12, 665145.	1.5	5
28	Prevalence of Osteoporosis and Fracture in China. JAMA Network Open, 2021, 4, e2121106.	2.8	178
29	Estimating the future clinical and economic benefits of improving osteoporosis diagnosis and treatment among women in China: a simulation projection model from 2020 to 2040. Archives of Osteoporosis, 2021, 16, 118.	1.0	13
30	Transcriptome landscape of the late-stage alcohol-induced osteonecrosis of the human femoral head. Bone, 2021, 150, 116012.	1.4	4
31	The Negative Impacts of Acromegaly on Bone Microstructure Not Fully Reversible. Frontiers in Endocrinology, 2021, 12, 738895.	1.5	9
32	Clinical, Biochemical, Radiological, Genetic and Therapeutic Analysis of Patients with COMP Gene Variants. Calcified Tissue International, 2021, , 1.	1.5	2
33	Atypical Femoral Fracture Associated With Overuse of Bisphosphonate Evaluated by High-Resolution Peripheral Quantitative Computed Tomography (HR-pQCT). Journal of Clinical Densitometry, 2020, 23, 329-334.	0.5	2
34	Nonremission and Recurrent Tumorâ€induced Osteomalacia: A Retrospective Study. Journal of Bone and Mineral Research, 2020, 35, 469-477.	3.1	43
35	Impaired geometry, volumetric density, and microstructure of cortical and trabecular bone assessed by HR-pQCT in both sporadic and MEN1-related primary hyperparathyroidism. Osteoporosis International, 2020, 31, 165-173.	1.3	14
36	Algorithm for the management of patients at low, high and very high risk of osteoporotic fractures. Osteoporosis International, 2020, 31, 1-12.	1.3	220

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37	Impaired bone microarchitecture in distal interphalangeal joints in patients with primary hypertrophic osteoarthropathy assessed by high-resolution peripheral quantitative computed tomography. Osteoporosis International, 2020, 31, 153-164.	1.3	6
38	Recurrent Femoral Fractures in a Boy with an Atypical Progeroid Syndrome: A Case Report. Calcified Tissue International, 2020, 106, 325-330.	1.5	2
39	Predicting the intervention threshold for initiating osteoporosis treatment among postmenopausal women in China: a cost-effectiveness analysis based on real-world data. Osteoporosis International, 2020, 31, 307-316.	1.3	18
40	Prevalence of Parathyroid Carcinoma and Atypical Parathyroid Neoplasms in 153 Patients With Multiple Endocrine Neoplasia Type 1: Case Series and Literature Review. Frontiers in Endocrinology, 2020, 11, 557050.	1.5	9
41	A novel mutation in PLS3 causes extremely rare Xâ€linked osteogenesis imperfecta. Molecular Genetics & Genomic Medicine, 2020, 8, e1525.	0.6	12
42	Pharmacologic intervention for prevention of fractures in osteopenic and osteoporotic postmenopausal women: Systemic review and meta-analysis. Bone Reports, 2020, 13, 100729.	0.2	14
43	Clinical Characteristics and Bone Features of Autosomal Recessive Hypophosphatemic Rickets Type 1 in Three Chinese Families: Report of Five Chinese Cases and Review of the Literature. Calcified Tissue International, 2020, 107, 636-648.	1.5	7
44	Bone mineral density and bone microarchitecture in a cohort of patients with Erdheim-Chester Disease. Orphanet Journal of Rare Diseases, 2020, 15, 236.	1.2	2
45	The first case report of Kyphoscoliotic Ehlers-Danlos syndrome of chinese origin with a novel PLOD1 gene mutation. BMC Medical Genetics, 2020, 21, 214.	2.1	3
46	Defective O-glycosylation of novel FGF23 mutations in a Chinese family with hyperphosphatemic familial tumoral calcinosis. Bone, 2020, 137, 115401.	1.4	3
47	Manifestations of left ventricular dysfunction and arrhythmia in patients with chronic hypoparathyroidism and pseudohypoparathyroidism: a preliminary study. BMC Endocrine Disorders, 2020, 20, 61.	0.9	3
48	Low HbA1c With Normal Hemoglobin in a Diabetes Patient Caused by PIEZO1 Gene Variant: A Case Report. Frontiers in Endocrinology, 2020, 11, 356.	1.5	4
49	Ageâ€, Siteâ€, and Sexâ€Specific Normative Centile Curves for <scp>HRâ€pQCT</scp>â€Derived Microarchitectural and Bone Strength Parameters in a Chinese Mainland Population. Journal of Bone and Mineral Research, 2020, 35, 2159-2170.	3.1	27
50	Germline GCM2 Mutation Screening in Chinese Primary Hyperparathyroidism Patients. Endocrine Practice, 2020, 26, 1093-1104.	1.1	9
51	Bone microstructure of adult patients with non-surgical hypoparathyroidism assessed by high-resolution peripheral quantitative computed tomography. Osteoporosis International, 2020, 31, 2219-2230.	1.3	6
52	A novel long-range deletion spanning STX16 and NPEPL1 causing imprinting defects of the GNAS locus discovered in a patient with autosomal-dominant pseudohypoparathyroidism type 1B. Endocrine, 2020, 69, 212-219.	1.1	14
53	Mutational landscape and genetic signatures of cellâ€free DNA in tumourâ€induced osteomalacia. Journal of Cellular and Molecular Medicine, 2020, 24, 4931-4943.	1.6	4
54	Effects of 24 Weeks of Growth Hormone Treatment on Bone Microstructure and Volumetric Bone Density in Patients with Childhood-Onset Adult GH Deficiency. International Journal of Endocrinology, 2020, 2020, 1-9.	0.6	5

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55	Early Discrimination Between Tumor-Induced Rickets/Osteomalacia and X-Linked Hypophosphatemia in Chinese Children and Adolescents: A Retrospective Caseâ€“Control Study. Journal of Bone and Mineral Research, 2020, 36, 1739-1748.	3.1	3
56	Alteration of Bone Density, Microarchitecture, and Strength in Patients with Camuratiâ€“Engelmann Disease: Assessed by HR-pQCT. Journal of Bone and Mineral Research, 2020, 37, 78-86.	3.1	4
57	Clinical and Genetic Findings in a Chinese Cohort of Patients with Digeorge Syndrome-Related Hypoparathyroidism. Endocrine Practice, 2020, 26, 642-650.	1.1	3
58	Bone Impairment in a Large Cohort of Chinese Patients With Tumor-Induced Osteomalacia Assessed by HR-pQCT and TBS. Journal of Bone and Mineral Research, 2020, 37, 454-464.	3.1	13
59	Bone Geometry, Density, Microstructure, and Biomechanical Properties in the Distal Tibia in Patients With Primary Hypertrophic Osteoarthropathy Assessed by Second-Generation High-Resolution Peripheral Quantitative Computed Tomography. Journal of Bone and Mineral Research, 2020, 37, 484-493.	3.1	4
60	East meets West: current practices and policies in the management of musculoskeletal aging. Aging Clinical and Experimental Research, 2019, 31, 1351-1373.	1.4	32
61	Genetic Screening in a Large Chinese Cohort of Childhood Onset Hypoparathyroidism by Next-Generation Sequencing Combined with TBX1-MLPA. Journal of Bone and Mineral Research, 2019, 34, 2254-2263.	3.1	21
62	Earlier Onset in Autosomal Dominant Hypophosphatemic Rickets of R179 than R176 Mutations in Fibroblast Growth Factor 23: Report of 20 Chinese Cases and Review of the Literature. Calcified Tissue International, 2019, 105, 476-486.	1.5	12
63	Incident Fracture Risk in Type 2 Diabetic Postmenopausal Women in Mainland China: Peking Vertebral Fracture Study. Calcified Tissue International, 2019, 105, 466-475.	1.5	16
64	Clinical and genetic analysis in a large Chinese cohort of patients with X-linked hypophosphatemia. Bone, 2019, 121, 212-220.	1.4	48
65	A 105Âkb interstitial insertion in the Xq27.1 palindrome from pseudoautosomal region PAR1 causes a novel X-linked recessive compound phenotype. Journal of Translational Medicine, 2019, 17, 138.	1.8	2
66	Eldecalcitol increases bone mineral density in Chinese osteoporotic patients without vitamin D or calcium supplementation. Journal of Bone and Mineral Metabolism, 2019, 37, 1036-1047.	1.3	14
67	The epidemiology of osteoporosis, associated fragility fractures, and management gap in China. Archives of Osteoporosis, 2019, 14, 32.	1.0	128
68	A novel missense mutation in P4HB causes mild osteogenesis imperfecta. Bioscience Reports, 2019, 39, .	1.1	12
69	Calciotropic Hormones and the Prevalence of Vertebral Fractures in Chinese Postmenopausal Women with Vitamin D Insufficiency: Peking Vertebral Fracture Study. Calcified Tissue International, 2019, 104, 622-630.	1.5	5
70	Health-related quality of life in children with osteogenesis imperfecta: a large-sample study. Osteoporosis International, 2019, 30, 461-468.	1.3	19
71	Mutation update and long-term outcome after treatment with active vitamin D3 in Chinese patients with pseudovitamin D-deficiency rickets (PDDR). Osteoporosis International, 2019, 30, 481-489.	1.3	8
72	Bone microarchitecture and volumetric bone density impairment in young male adults with childhood-onset growth hormone deficiency. European Journal of Endocrinology, 2019, 180, 145-153.	1.9	20

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73	The first case of primary hypertrophic osteoarthropathy with soft tissue giant tumors caused by HPGD loss-of-function mutation. Endocrine Connections, 2019, 8, 736-744.	0.8	5
74	Novel compound heterozygous mutations in SERPINH1 cause rare autosomal recessive osteogenesis imperfecta type X. Osteoporosis International, 2018, 29, 1389-1396.	1.3	13
75	Effects of alendronate and alfacalcidol on bone in patients with myasthenia gravis initiating glucocorticoids treatment. Clinical Endocrinology, 2018, 88, 380-387.	1.2	5
76	Four novel mutations in the ALPL gene in Chinese patients with odonto, childhood, and adult hypophosphatasia. Bioscience Reports, 2018, 38, .	1.1	11
77	An updated hip fracture projection in Asia: The Asian Federation of Osteoporosis Societies study. Osteoporosis and Sarcopenia, 2018, 4, 16-21.	0.7	222
78	Identification of mutations in the prostaglandin transporter gene SLCO2A1 and phenotypic comparison between two subtypes of primary hypertrophic osteoarthropathy (PHO): A single-center study. Bone, 2018, 106, 96-102.	1.4	27
79	Novel Mutations in PLOD2 Cause Rare Bruck Syndrome. Calcified Tissue International, 2018, 102, 296-309.	1.5	22
80	Clinical Management of Malignant Insulinoma: a single Institutionâ€™s experience over three decades. BMC Endocrine Disorders, 2018, 18, 92.	0.9	20
81	Tumor-induced osteomalacia. Osteoporosis and Sarcopenia, 2018, 4, 119-127.	0.7	40
82	Efficacy and safety of alfacalcidol in Chinese postmenopausal women aged over 65 with osteoporosis or osteopenia. Medicine (United States), 2018, 97, e13159.	0.4	4
83	URAT1 mutations cause renal hypouricaemia combined with Fanconi syndrome in a Chinese patient. Nephrology, 2018, 23, 797-798.	0.7	0
84	Consensus on best practice standards for Fracture Liaison Service in the Asia-Pacific region. Archives of Osteoporosis, 2018, 13, 59.	1.0	14
85	Vertebral fractures among breast cancer survivors in China: a cross-sectional study of prevalence and health services gaps. BMC Cancer, 2018, 18, 104.	1.1	6
86	Assessment of bone quality in patients with diabetes mellitus. Osteoporosis International, 2018, 29, 1721-1736.	1.3	55
87	Prevalence of Vitamin D Inadequacy Among Chinese Postmenopausal Women: A Nationwide, Multicenter, Cross-Sectional Study. Frontiers in Endocrinology, 2018, 9, 782.	1.5	16
88	Pharmacological Treatment of Bone Loss. Current Pharmaceutical Design, 2018, 23, 6298-6301.	0.9	3
89	Association between bone mineral density, muscle strength, and vitamin D status in patients with myasthenia gravis: a cross-sectional study. Osteoporosis International, 2017, 28, 2383-2390.	1.3	9
90	A novel large fragment deletion in PLS3 causes rare X-linked early-onset osteoporosis and response to zoledronic acid. Osteoporosis International, 2017, 28, 2691-2700.	1.3	24

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91	Vertebral fracture in postmenopausal Chinese women: a population-based study. Osteoporosis International, 2017, 28, 2583-2590.	1.3	47
92	Familial Early-Onset Pagetâ€™s Disease of Bone Associated with a Novel hnRNPA2B1 Mutation. Calcified Tissue International, 2017, 101, 159-169.	1.5	35
93	Associations between FGF21, osteonectin and bone turnover markers in type 2 diabetic patients with albuminuria. Journal of Diabetes and Its Complications, 2017, 31, 583-588.	1.2	10
94	Primary hyperparathyroidism in Chinese children and adolescents: A singleâ€centre experience at Peking Union Medical College Hospital. Clinical Endocrinology, 2017, 87, 865-873.	1.2	20
95	Atypical skeletal manifestations of rickets in a familial hypocalciuric hypercalcemia patient. Bone Research, 2017, 5, 17001.	5.4	6
96	Gene mutation spectrum and genotype-phenotype correlation in a cohort of Chinese osteogenesis imperfecta patients revealed by targeted next generation sequencing. Osteoporosis International, 2017, 28, 2985-2995.	1.3	56
97	The Asian Federation of Osteoporosis Societies' call to action to improve the undertreatment of osteoporosis in Asia. Osteoporosis and Sarcopenia, 2017, 3, 161-163.	0.7	10
98	Mild Camurati-Engelamann disease presenting with exophthalmos as the first and only manifestation: A case report. Molecular Medicine Reports, 2016, 14, 2710-2716.	1.1	2
99	Clinical and Genetic Analysis of Multiple Endocrine Neoplasia Type 1-Related Primary Hyperparathyroidism in Chinese. PLoS ONE, 2016, 11, e0166634.	1.1	18
100	Clinical and genetic findings in a Chinese family with VDR-associated hereditary vitamin D-resistant rickets. Bone Research, 2016, 4, 16018.	5.4	7
101	Novel COL2A1 mutations causing spondyloepiphyseal dysplasia congenita in three unrelated Chinese families. European Spine Journal, 2016, 25, 2967-2974.	1.0	7
102	Osteogenesis imperfecta type V: Genetic and clinical findings in eleven Chinese patients. Clinica Chimica Acta, 2016, 462, 201-209.	0.5	10
103	Genotype-phenotype analysis of a rare type of osteogenesis imperfecta in four Chinese families with WNT1 mutations. Clinica Chimica Acta, 2016, 461, 172-180.	0.5	21
104	Identification of suitable reference gene and biomarkers of serum miRNAs for osteoporosis. Scientific Reports, 2016, 6, 36347.	1.6	85
105	Establishment of a normal reference value of parathyroid hormone in a large healthy Chinese population and evaluation of its relation to bone turnover and bone mineral density. Osteoporosis International, 2016, 27, 1907-1916.	1.3	45
106	Novel mutations of CLCN7 cause autosomal dominant osteopetrosis type II (ADO-II) and intermediate autosomal recessive osteopetrosis (IARO) in Chinese patients. Osteoporosis International, 2016, 27, 1047-1055.	1.3	23
107	MEN1 c.825-1G>A mutation in a family with multiple endocrine neoplasia type 1: A case report. Molecular Medicine Reports, 2015, 12, 6152-6156.	1.1	3
108	Chimeric CYP11B2 / CYP11B1 causing 11Î²-hydroxylase deficiency in Chinese patients with congenital adrenal hyperplasia. Steroids, 2015, 101, 51-55.	0.8	12

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109	Surgical Treatments of Tumor-Induced Osteomalacia Lesions in Long Bones. Journal of Bone and Joint Surgery - Series A, 2015, 97, 1084-1094.	1.4	29
110	A survey of outcomes and management of patients post fragility fractures in China. Osteoporosis International, 2015, 26, 2631-2640.	1.3	44
111	Novel WISP3 mutations causing spondyloepiphyseal dysplasia tarda with progressive arthropathy in two unrelated Chinese families. Joint Bone Spine, 2015, 82, 125-128.	0.8	14
112	Aromatase deficiency in a Chinese adult man caused by novel compound heterozygous CYP19A1 mutations: Effects of estrogen replacement therapy on the bone, lipid, liver and glucose metabolism. Molecular and Cellular Endocrinology, 2015, 399, 32-42.	1.6	46
113	Management of osteoporosis with calcitriol in elderly Chinese patients: a systematic review. Clinical Interventions in Aging, 2014, 9, 515.	1.3	10
114	The First Mutation Identified in a Chinese Acrodysostosis Patient Confirms a p.G289E Variation of PRKAR1A Causes Acrodysostosis. International Journal of Molecular Sciences, 2014, 15, 13267-13274.	1.8	8
115	Association of GALNT3 gene polymorphisms with bone mineral density in Chinese postmenopausal women. Menopause, 2014, 21, 515-521.	0.8	9
116	Suppressed bone turnover was associated with increased osteoporotic fracture risks in non-obese postmenopausal Chinese women with type 2 diabetes mellitus. Osteoporosis International, 2014, 25, 1999-2005.	1.3	70
117	A compound heterozygous mutation in SLC34A3 causes hereditary hypophosphatemic rickets with hypercalciuria in a Chinese patient. Bone, 2014, 59, 114-121.	1.4	17
118	A4.14â€…Increased bone turnover after switch to tenofovir + lopinavir/ritonavir in chinese HIV + patients. Annals of the Rheumatic Diseases, 2014, 73, A62.1-A62.	0.5	0
119	A4.15â€…Osteoporosis knowledge, self-efficacy, and health beliefs among chinese men and women with HIV. Annals of the Rheumatic Diseases, 2014, 73, A62.2-A63.	0.5	0
120	Chinese Bone Turnover Marker Study: Reference Ranges for C-Terminal Telopeptide of Type I Collagen and Procollagen I N-Terminal Peptide by Age and Gender. PLoS ONE, 2014, 9, e103841.	1.1	43
121	Novel Mutations in FKBP10 and PLOD2 Cause Rare Bruck Syndrome in Chinese Patients. PLoS ONE, 2014, 9, e107594.	1.1	26
122	Association of farnesyl diphosphate synthase polymorphisms and response to alendronate treatment in Chinese postmenopausal women with osteoporosis. Chinese Medical Journal, 2014, 127, 662-8.	0.9	6
123	FGF23 and Phosphate Wasting Disorders. Bone Research, 2013, 1, 120-132.	5.4	54
124	A haplotype of MATN3 is associated with vertebral fracture in Chinese postmenopausal women: Peking Vertebral Fracture (PK-VF) study. Bone, 2012, 50, 917-924.	1.4	8
125	Novel mutations of CYP27B1 gene lead to reduced activity of 1Î±-hydroxylase in Chinese patients. Bone, 2012, 51, 563-569.	1.4	25
126	Exome Sequencing Identifies SLCO2A1 Mutations as a Cause of Primary Hypertrophic Osteoarthropathy. American Journal of Human Genetics, 2012, 90, 125-132.	2.6	157

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127	FGF23 analysis of a Chinese family with autosomal dominant hypophosphatemic rickets. Journal of Bone and Mineral Metabolism, 2012, 30, 78-84.	1.3	16
128	The levels of bone turnover markers in Chinese postmenopausal women. Menopause, 2011, 18, 1237-1243.	0.8	65
129	Three Novel Mutations of the PHEX Gene in Three Chinese Families with X-linked Dominant Hypophosphatemic Rickets. Calcified Tissue International, 2007, 81, 415-420.	1.5	25
130	Association of genetic variants of vit D binding protein (DBP/GC) and of the enzyme catalyzing its 25-hydroxylation (DCYP2R1) and serum vit D in postmenopausal women. Hormones, 2002, 13, 345-52.	0.9	14
131	Bone Volumetric Density, Microarchitecture, and Estimated Bone Strength in Tumor-Induced Rickets/Osteomalacia Versus X-linked Hypophosphatemia in Chinese Adolescents. Frontiers in Endocrinology, 0, 13, .	1.5	2
132	Na-Cl Co-transporter (NCC) gene inactivation is associated with improved bone microstructure. Osteoporosis International, 0, , .	1.3	0
133	A Novel Synonymous Variant of PHEX in a Patient with X-Linked Hypophosphatemia. Calcified Tissue International, 0, , .	1.5	1