Frank Rauch

List of Publications by Year in Descending Order

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

118 366 17,557 72 h-index g-index citations papers 6.69 19,820 387 4.7 avg, IF L-index ext. citations ext. papers

#	Paper	IF	Citations
366	Maternal excess adiposity and serum 25-hydroxyvitamin D . <i>BMC Pregnancy and Childbirth</i> , 2022 , 22, 83	3.2	0
365	Characterization and functional analysis of the adipose tissue-derived stromal vascular fraction of pediatric patients with osteogenesis imperfecta <i>Scientific Reports</i> , 2022 , 12, 2414	4.9	1
364	Bone Turnover Markers in Children: From Laboratory Challenges to Clinical Interpretation <i>Calcified Tissue International</i> , 2022 , 1	3.9	O
363	3D image registration marginally improves the precision of HR-pQCT measurements compared to cross-sectional-area registration in adults with osteogenesis imperfecta <i>Journal of Bone and Mineral Research</i> , 2022 ,	6.3	1
362	Dominant osteogenesis imperfecta with low bone turnover caused by a heterozygous SP7 variant <i>Bone</i> , 2022 , 116400	4.7	1
361	Correction of neonatal vitamin D status using 1000 IU/d of vitamin D increased lean body mass by 12 months of age compared to 400 IU/d: a rand1omized controlled trial <i>American Journal of Clinical Nutrition</i> , 2022 ,	7	2
360	Maternal Vitamin D Status and Gestational Weight Gain as Correlates of Neonatal Bone Mass in Healthy Term Breastfed Young Infants from Montreal, Canada <i>Nutrients</i> , 2021 , 13,	6.7	2
359	A Validated Risk Prediction Model for Bone Fragility in Children With Acute Lymphoblastic Leukemia. <i>Journal of Bone and Mineral Research</i> , 2021 , 36, 2290	6.3	1
358	Histomorphometric and microarchitectural analysis of bone in metastatic breast cancer patients. <i>Bone Reports</i> , 2021 , 15, 101145	2.6	
357	A standard set of outcome measures for the comprehensive assessment of osteogenesis imperfecta. <i>Orphanet Journal of Rare Diseases</i> , 2021 , 16, 140	4.2	3
356	Increased Osteocyte Lacunae Density in the Hypermineralized Bone Matrix of Children with Osteogenesis Imperfecta Type I. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	2
355	The Accuracy of Incident Vertebral Fracture Detection in Children Using Targeted Case-Finding Approaches. <i>Journal of Bone and Mineral Research</i> , 2021 , 36, 1255-1268	6.3	0
354	Compressive Strength of Iliac Bone ECM Is Not Reduced in Osteogenesis Imperfecta and Increases With Mineralization. <i>Journal of Bone and Mineral Research</i> , 2021 , 36, 1364-1375	6.3	4
353	Effects of treatment with a bone-targeted prostaglandin E2 receptor 4 agonist C3 (Mes-1007) in a mouse model of severe osteogenesis imperfecta. <i>Bone</i> , 2021 , 145, 115867	4.7	O
352	Reply to: Burosumab for Tumor-Induced Osteomalacia: not Enough of a Good Thing. <i>Journal of Bone and Mineral Research</i> , 2021 ,	6.3	1
351	Calvaria Bone Transcriptome in Mouse Models of Osteogenesis Imperfecta. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	1
350	Multisite longitudinal calibration of HR-pQCT scanners and precision in osteogenesis imperfecta. <i>Bone</i> , 2021 , 147, 115880	4.7	1

349	Osteogenesis imperfecta tooth level phenotype analysis: Cross-sectional study. <i>Bone</i> , 2021 , 147, 11591	7 4.7	1
348	Male but not female mice with severe osteogenesis imperfecta are partially protected from high-fat diet-induced obesity. <i>Molecular Genetics and Metabolism</i> , 2021 , 133, 211-221	3.7	2
347	Insufficient Vitamin D Status at Birth Is Corrected by Vitamin D Supplementation (1000 IU/Day) With Increases in Lean Mass Evident at 12 Months of Age in Healthy Term Infants. <i>Current Developments in Nutrition</i> , 2021 , 5, 804-804	0.4	78
346	Osteoporotic Fractures and Vertebral Body Reshaping in Children With Glucocorticoid-treated Rheumatic Disorders. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021 , 106, e5195-e5207	5.6	1
345	Predicting ambulatory function at skeletal maturity in children with moderate to severe osteogenesis imperfecta. <i>European Journal of Pediatrics</i> , 2021 , 180, 233-239	4.1	2
344	Burosumab for the Treatment of Tumor-Induced Osteomalacia. <i>Journal of Bone and Mineral Research</i> , 2021 , 36, 627-635	6.3	29
343	Prevalence of SARS-CoV-2 infections in a pediatric orthopedic hospital. <i>Paediatric Anaesthesia</i> , 2021 , 31, 247-248	1.8	1
342	Health-related quality of life in adults with osteogenesis imperfecta. Clinical Genetics, 2021, 99, 772-779	94	Ο
341	Muscle transcriptome in mouse models of osteogenesis imperfecta. <i>Bone</i> , 2021 , 148, 115940	4.7	2
340	Experiences of Children With Osteogenesis Imperfecta in the Co-design of the Interactive Assessment and Communication Tool Sisom OI: Secondary Analysis of Qualitative Design Sessions. JMIR Pediatrics and Parenting, 2021, 4, e22784	4.2	O
339	Craniocervical abnormalities in osteogenesis imperfecta type V. <i>Osteoporosis International</i> , 2021 , 33, 177	5.3	0
338	Missing and unerupted teeth in osteogenesis imperfecta. <i>Bone</i> , 2021 , 150, 116011	4.7	O
337	Muscle-bone properties after prolonged voluntary wheel running in a mouse model of dominant severe osteogenesis imperfecta. <i>Journal of Musculoskeletal Neuronal Interactions</i> , 2021 , 21, 517-527	1.3	
336	Bone Disease in Patients with Ehlers-Danlos Syndromes. <i>Current Osteoporosis Reports</i> , 2020 , 18, 95-102	5.4	11
335	Malocclusion traits and oral health-related quality of life in children with osteogenesis imperfecta: A cross-sectional study. <i>Journal of the American Dental Association</i> , 2020 , 151, 480-490.e2	1.9	1
334	Treatment response to long term antiresorptive therapy in osteogenesis imperfecta type VI: does genotype matter?. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2020 , 33, 1617-1624	1.6	1
333	Giving Children With Osteogenesis Imperfecta a Voice: Participatory Approach for the Development of the Interactive Assessment and Communication Tool Sisom OI. <i>Journal of Medical Internet Research</i> , 2020 , 22, e17947	7.6	1
332	Bone densities and bone geometry in children and adolescents with arthrogryposis. <i>Bone</i> , 2020 , 137, 115454	4.7	1

331	Increased Burden of Common Risk Alleles in Children With a Significant Fracture History. <i>Journal of Bone and Mineral Research</i> , 2020 , 35, 875-882	6.3	4
330	The Accuracy of Prevalent Vertebral Fracture Detection in Children Using Targeted Case-Finding Approaches. <i>Journal of Bone and Mineral Research</i> , 2020 , 35, 460-468	6.3	7
329	Bladder and bowel symptoms experienced by children with osteogenesis imperfecta. <i>Jornal De Pediatria (Versi</i>) Em Portuguis), 2020 , 96, 472-478	0.2	
328	Assessment of longitudinal bone growth in osteogenesis imperfecta using metacarpophalangeal pattern profiles. <i>Bone</i> , 2020 , 140, 115547	4.7	2
327	Musculoskeletal phenotype in two unrelated individuals with a recurrent nonsense variant in SGMS2. <i>Bone</i> , 2020 , 134, 115261	4.7	9
326	Exploring the Perceived Self-management Needs of Young Adults With Osteogenesis Imperfecta. <i>Clinical Nurse Specialist</i> , 2020 , 34, 99-106	0.6	1
325	Bladder and bowel symptoms experienced by children with osteogenesis imperfecta. <i>Jornal De Pediatria</i> , 2020 , 96, 472-478	2.6	O
324	A de novo frameshift FGFR1 mutation extending the protein in an individual with multiple epiphyseal dysplasia and hypogonadotropic hypogonadism without anosmia. <i>European Journal of Medical Genetics</i> , 2020 , 63, 103784	2.6	2
323	Mineralized tissues in hypophosphatemic rickets. <i>Pediatric Nephrology</i> , 2020 , 35, 1843-1854	3.2	12
322	Validation of the Adolescent Pediatric Pain Tool for the Multidimensional Measurement of Pain in Children and Adolescents Diagnosed with Osteogenesis Imperfecta <i>Canadian Journal of Pain</i> , 2019 , 3, 148-156	1.5	O
321	Osteogenesis Imperfecta: New Perspectives From Clinical and Translational Research. <i>JBMR Plus</i> , 2019 , 3, e10174	3.9	45
320	Identification of genetic variants associated with skeletal muscle function deficit in childhood acute lymphoblastic leukemia survivors. <i>Pharmacogenomics and Personalized Medicine</i> , 2019 , 12, 33-45	2.1	2
319	A Multicenter Observational Cohort Study to Evaluate the Effects of Bisphosphonate Exposure on Bone Mineral Density and Other Health Outcomes in Osteogenesis Imperfecta. <i>JBMR Plus</i> , 2019 , 3, e10	13:8	7
318	Development of the Good2Go MyHealth Passport for individuals with Osteogenesis Imperfecta: A knowledge-synthesis study. <i>International Journal of Orthopaedic and Trauma Nursing</i> , 2019 , 33, 27-34	1.1	4
317	Management of bone disease in cystinosis: Statement from an international conference. <i>Journal of Inherited Metabolic Disease</i> , 2019 , 42, 1019-1029	5.4	28
316	Identification of a single-nucleotide polymorphism within gene associated with bone morbidity in childhood acute lymphoblastic leukemia survivors. <i>Pharmacogenomics</i> , 2019 , 20, 409-420	2.6	3
315	The Bone Phenotype and Pain Response to Pamidronate in Tyrosine Kinase Inhibitor-Treated Chronic Myelogenous Leukemia. <i>Journal of the Endocrine Society</i> , 2019 , 3, 857-864	0.4	5
314	Mendelian bone fragility disorders. <i>Bone</i> , 2019 , 126, 11-17	4.7	14

(2018-2019)

313	Mobility in osteogenesis imperfecta: a multicenter North American study. <i>Genetics in Medicine</i> , 2019 , 21, 2311-2318	8.1	6
312	ALPL mutations in adults with rheumatologic disorders and low serum alkaline phosphatase activity. <i>Journal of Bone and Mineral Metabolism</i> , 2019 , 37, 893-899	2.9	6
311	Growth characteristics in individuals with osteogenesis imperfecta in North America: results from a multicenter study. <i>Genetics in Medicine</i> , 2019 , 21, 275-283	8.1	15
310	Novel ActRIIB ligand trap increases muscle mass and improves bone geometry in a mouse model of severe osteogenesis imperfecta. <i>Bone</i> , 2019 , 128, 115036	4.7	14
309	Burosumab Improved Histomorphometric Measures of Osteomalacia in Adults with X-Linked Hypophosphatemia: A Phase 3, Single-Arm, International Trial. <i>Journal of Bone and Mineral Research</i> , 2019 , 34, 2183-2191	6.3	41
308	Osteogenesis Imperfecta: Skeletal Outcomes After Bisphosphonate Discontinuation at Final Height. <i>Journal of Bone and Mineral Research</i> , 2019 , 34, 2198-2204	6.3	3
307	Caries prevalence and experience in individuals with osteogenesis imperfecta: A cross-sectional multicenter study. <i>Special Care in Dentistry</i> , 2019 , 39, 214-219	1.7	6
306	Five-year efficacy and safety of asfotase alfa therapy for adults and adolescents with hypophosphatasia. <i>Bone</i> , 2019 , 121, 149-162	4.7	55
305	Cone-Beam Computed Tomography of Osteogenesis Imperfecta Types III and IV: Three-Dimensional Evaluation of Craniofacial Features and Upper Airways. <i>JBMR Plus</i> , 2019 , 3, e10124	3.9	3
304	Impact of Vertebral Fractures and Glucocorticoid Exposure on Height Deficits in Children During Treatment of Leukemia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 , 104, 213-222	5.6	6
303	Oro-dental and cranio-facial characteristics of osteogenesis imperfecta type V. <i>European Journal of Medical Genetics</i> , 2019 , 62, 103606	2.6	3
302	Effect of Anti-TGF-Treatment in a Mouse Model of Severe Osteogenesis Imperfecta. <i>Journal of Bone and Mineral Research</i> , 2019 , 34, 207-214	6.3	25
301	Dental and craniofacial characteristics caused by the p.Ser40Leu mutation in IFITM5. <i>American Journal of Medical Genetics, Part A</i> , 2019 , 179, 65-70	2.5	5
300	Vitamin D nutritional status and bone turnover markers in childhood acute lymphoblastic leukemia survivors: A PETALE study. <i>Clinical Nutrition</i> , 2019 , 38, 912-919	5.9	11
299	Whole-Exome Sequencing Identifies an Intronic Cryptic Splice Site in Causing Osteogenesis Imperfecta Type VI. <i>JBMR Plus</i> , 2018 , 2, 235-239	3.9	2
298	Pain and quality of life of children and adolescents with osteogenesis imperfecta over a bisphosphonate treatment cycle. <i>European Journal of Pediatrics</i> , 2018 , 177, 891-902	4.1	11
297	Genotype and malocclusion in patients with osteogenesis imperfecta. <i>Orthodontics and Craniofacial Research</i> , 2018 , 21, 71-77	3	3
296	Anabolic Therapy for the Treatment of Osteoporosis in Childhood. <i>Current Osteoporosis Reports</i> , 2018 , 16, 269-276	5.4	16

295	Pulmonary and diaphragmatic pathology in collagen type I II mutant mice with osteogenesis imperfecta. <i>Pediatric Research</i> , 2018 , 83, 1165-1171	3.2	13
294	Vitamin D status and functional health outcomes in children aged 2-8 y: a 6-mo vitamin D randomized controlled trial. <i>American Journal of Clinical Nutrition</i> , 2018 , 107, 355-364	7	11
293	Assessment of the effect of systemic delivery of sclerostin antibodies on Wnt signaling in distraction osteogenesis. <i>Journal of Bone and Mineral Metabolism</i> , 2018 , 36, 373-382	2.9	4
292	Novel WNT1 mutations in children with osteogenesis imperfecta: Clinical and functional characterization. <i>Bone</i> , 2018 , 114, 144-149	4.7	20
291	Hypercalcemia and hypercalciuria during denosumab treatment in children with osteogenesis imperfecta type VI. <i>Journal of Musculoskeletal Neuronal Interactions</i> , 2018 , 18, 76-80	1.3	27
290	Crispr-Cas9 engineered osteogenesis imperfecta type V leads to severe skeletal deformities and perinatal lethality in mice. <i>Bone</i> , 2018 , 107, 131-142	4.7	24
289	A Best Practice Initiative to Optimize Transfer of Young Adults With Osteogenesis Imperfecta From Child to Adult Healthcare Services. <i>Clinical Nurse Specialist</i> , 2018 , 32, 323-335	0.6	3
288	Oral health-related quality of life in children and adolescents with osteogenesis imperfecta: cross-sectional study. <i>Orphanet Journal of Rare Diseases</i> , 2018 , 13, 187	4.2	10
287	A multicenter study to evaluate pulmonary function in osteogenesis imperfecta. <i>Clinical Genetics</i> , 2018 , 94, 502-511	4	20
286	Osteogenesis imperfecta type III/Ehlers-Danlos overlap syndrome in a Chinese man. <i>Intractable and Rare Diseases Research</i> , 2018 , 7, 37-41	1.4	6
285	Bone Morbidity and Recovery in Children With Acute Lymphoblastic Leukemia: Results of a Six-Year Prospective Cohort Study. <i>Journal of Bone and Mineral Research</i> , 2018 , 33, 1435-1443	6.3	50
284	Hip Dysplasia in Children With Osteogenesis Imperfecta: Association With Collagen Type I C-Propeptide Mutations. <i>Journal of Pediatric Orthopaedics</i> , 2017 , 37, 479-483	2.4	10
283	Static Postural Control in Youth With Osteogenesis Imperfecta Type I. <i>Archives of Physical Medicine and Rehabilitation</i> , 2017 , 98, 1948-1954	2.8	3
282	Muscle Function in Osteogenesis Imperfecta Type IV. Calcified Tissue International, 2017, 101, 362-370	3.9	14
281	Learning from the experience of a long-standing interprofessional osteogenesis imperfecta clinic: A case study evaluation. <i>Journal of Interprofessional Education and Practice</i> , 2017 , 7, 54-60	0.6	
280	Hypermineralization and High Osteocyte Lacunar Density in Osteogenesis Imperfecta Type V Bone Indicate Exuberant Primary Bone Formation. <i>Journal of Bone and Mineral Research</i> , 2017 , 32, 1884-1892	6.3	34
279	Molecular diagnosis in children with fractures but no extraskeletal signs of osteogenesis imperfecta. <i>Osteoporosis International</i> , 2017 , 28, 2095-2101	5.3	21
278	The PETALE study: Late adverse effects and biomarkers in childhood acute lymphoblastic leukemia survivors. <i>Pediatric Blood and Cancer</i> , 2017 , 64, e26361	3	43

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277	Diaphyseal Femur Fractures in Osteogenesis Imperfecta: Characteristics and Relationship With Bisphosphonate Treatment. <i>Journal of Bone and Mineral Research</i> , 2017 , 32, 1034-1039	6.3	24
276	DHA and EPA in red blood cell membranes are associated with dietary intakes of omega-3-rich fish in healthy children. <i>Prostaglandins Leukotrienes and Essential Fatty Acids</i> , 2017 , 124, 11-16	2.8	15
275	Muscle-Bone Interactions in Pediatric Bone Diseases. <i>Current Osteoporosis Reports</i> , 2017 , 15, 425-432	5.4	19
274	Long-term follow-up in osteogenesis imperfecta type VI. Osteoporosis International, 2017, 28, 2975-298	33 _{5.3}	12
273	Metabolic phenotype in the mouse model of osteogenesis imperfecta. <i>Journal of Endocrinology</i> , 2017 , 234, 279-289	4.7	14
272	Increased bone matrix mineralization in treatment-nalle children with inflammatory bowel disease. <i>Bone</i> , 2017 , 105, 50-56	4.7	8
271	An Official American Thoracic Society Workshop Report: Translational Research in Rare Respiratory Diseases. <i>Annals of the American Thoracic Society</i> , 2017 , 14, 1239-1247	4.7	1
270	Musculoskeletal health in newly diagnosed children with CrohnB disease. <i>Osteoporosis International</i> , 2017 , 28, 3169-3177	5.3	20
269	The Determinants of Peak Bone Mass. <i>Journal of Pediatrics</i> , 2017 , 180, 261-269	3.6	99
268	Geometry reconstruction method for patient-specific finite element models for the assessment of tibia fracture risk in osteogenesis imperfecta. <i>Medical and Biological Engineering and Computing</i> , 2017 , 55, 549-560	3.1	8
267	Bone Structural Characteristics and Response to Bisphosphonate Treatment in Children With Hajdu-Cheney Syndrome. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017 , 102, 4163-4172	5.6	18
266	Spondyloepimetaphysial Dysplasia with Joint Laxity in Three Siblings with Mutations. <i>Molecular Syndromology</i> , 2017 , 8, 303-307	1.5	5
265	The brains of the bones: how osteocytes use WNT1 to control bone formation. <i>Journal of Clinical Investigation</i> , 2017 , 127, 2539-2540	15.9	6
264	Muscle abnormalities in osteogenesis imperfecta. <i>Journal of Musculoskeletal Neuronal Interactions</i> , 2017 , 17, 1-7	1.3	13
263	Cesarean delivery is not associated with decreased at-birth fracture rates in osteogenesis imperfecta. <i>Genetics in Medicine</i> , 2016 , 18, 570-6	8.1	27
262	DNA sequence analysis in 598 individuals with a clinical diagnosis of osteogenesis imperfecta: diagnostic yield and mutation spectrum. <i>Osteoporosis International</i> , 2016 , 27, 3607-3613	5.3	85
261	Non-Lethal Type VIII Osteogenesis Imperfecta Has Elevated Bone Matrix Mineralization. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016 , 101, 3516-25	5.6	18
260	Sclerostin: More than a bone formation brake. <i>Science Translational Medicine</i> , 2016 , 8, 330fs7	17.5	3

259	The Spectrum of Recovery From Fracture-Induced Vertebral Deformity in Pediatric Leukemia. <i>Pediatric Blood and Cancer</i> , 2016 , 63, 1107-10	3	15
258	Histomorphometry and Bone Matrix Mineralization Before and After Bisphosphonate Treatment in Boys With Duchenne Muscular Dystrophy: A Paired Transiliac Biopsy Study. <i>Journal of Bone and Mineral Research</i> , 2016 , 31, 1060-9	6.3	31
257	Does Sclerostin Depletion Stimulate Fracture Healing in a Mouse Model?. <i>Clinical Orthopaedics and Related Research</i> , 2016 , 474, 1294-302	2.2	16
256	Body Composition in Children and Adolescents with Osteogenesis Imperfecta. <i>Journal of Pediatrics</i> , 2016 , 169, 232-7	3.6	20
255	Craniofacial and Dental Defects in the Col1a1Jrt/+ Mouse Model of Osteogenesis Imperfecta. Journal of Dental Research, 2016 , 95, 761-8	8.1	18
254	Bone matrix hypermineralization in prolyl-3 hydroxylase 1 deficient mice. <i>Bone</i> , 2016 , 85, 15-22	4.7	10
253	Osteogenesis Imperfecta Type VI in Individuals from Northern Canada. <i>Calcified Tissue International</i> , 2016 , 98, 566-72	3.9	24
252	The psychosocial experience of individuals living with osteogenesis imperfecta: a mixed-methods systematic review. <i>Quality of Life Research</i> , 2016 , 25, 1877-96	3.7	33
251	Effect of high-dose vitamin D supplementation on bone density in youth with osteogenesis imperfecta: A randomized controlled trial. <i>Bone</i> , 2016 , 86, 36-42	4.7	20
250	Scoliosis in osteogenesis imperfecta caused by COL1A1/COL1A2 mutations - genotype-phenotype correlations and effect of bisphosphonate treatment. <i>Bone</i> , 2016 , 86, 53-7	4.7	43
249	Dietary vitamin D dose-response in healthy children 2 to 8 y of age: a 12-wk randomized controlled trial using fortified foods. <i>American Journal of Clinical Nutrition</i> , 2016 , 103, 144-52	7	25
248	Agreement of spatio-temporal gait parameters between a vertical ground reaction force decomposition algorithm and a motion capture system. <i>Gait and Posture</i> , 2016 , 43, 257-64	2.6	12
247	Evaluation of a Modified Pamidronate Protocol for the Treatment of Osteogenesis Imperfecta. <i>Calcified Tissue International</i> , 2016 , 98, 42-8	3.9	7
246	Osteogenesis Imperfecta Type I Caused by COL1A1 Deletions. <i>Calcified Tissue International</i> , 2016 , 98, 76-84	3.9	22
245	Involving Families with Osteogenesis Imperfecta in Health Service Research: Joint Development of the OI/ECE Questionnaire. <i>PLoS ONE</i> , 2016 , 11, e0147654	3.7	5
244	Pediatric data for dual X-ray absorptiometric measures of normal lumbar bone mineral density in children under 5 years of age using the lunar prodigy densitometer. <i>Journal of Musculoskeletal Neuronal Interactions</i> , 2016 , 16, 247-55	1.3	4
243	Aging Versus Postmenopausal Osteoporosis: Bone Composition and Maturation Kinetics at Actively-Forming Trabecular Surfaces of Female Subjects Aged 1 to 84 Years. <i>Journal of Bone and Mineral Research</i> , 2016 , 31, 347-57	6.3	45
242	Evidence for a Role for Nanoporosity and Pyridinoline Content in Human Mild Osteogenesis Imperfecta. <i>Journal of Bone and Mineral Research</i> , 2016 , 31, 1050-9	6.3	29

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241	Osteogenesis imperfecta in children and adolescents-new developments in diagnosis and treatment. <i>Osteoporosis International</i> , 2016 , 27, 3427-3437	5.3	102
240	Quality of life in osteogenesis imperfecta: A mixed-methods systematic review. <i>American Journal of Medical Genetics, Part A</i> , 2016 , 170A, 62-76	2.5	37
239	Multidisciplinary Treatment of Severe Osteogenesis Imperfecta: Functional Outcomes at Skeletal Maturity. <i>Archives of Physical Medicine and Rehabilitation</i> , 2015 , 96, 1834-9	2.8	31
238	The effect of SERPINF1 in-frame mutations in osteogenesis imperfecta type VI. <i>Bone</i> , 2015 , 76, 115-20	4.7	15
237	A cross-sectional multicenter study of osteogenesis imperfecta in North America - results from the linked clinical research centers. <i>Clinical Genetics</i> , 2015 , 87, 133-40	4	45
236	The choice of normative pediatric reference database changes spine bone mineral density Z-scores but not the relationship between bone mineral density and prevalent vertebral fractures. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015 , 100, 1018-27	5.6	45
235	Behavioral signs of pain and functional impairment in a mouse model of osteogenesis imperfecta. <i>Bone</i> , 2015 , 81, 400-406	4.7	20
234	Incident Vertebral Fractures in Children With Leukemia During the Four Years Following Diagnosis. Journal of Clinical Endocrinology and Metabolism, 2015 , 100, 3408-17	5.6	68
233	A polyadenylation site variant causes transcript-specific BMP1 deficiency and frequent fractures in children. <i>Human Molecular Genetics</i> , 2015 , 24, 516-24	5.6	32
232	Osteotomy Healing in Children With Osteogenesis Imperfecta Receiving Bisphosphonate Treatment. <i>Journal of Bone and Mineral Research</i> , 2015 , 30, 1362-8	6.3	44
231	Incident Vertebral Fractures and Risk Factors in the First Three Years Following Glucocorticoid Initiation Among Pediatric Patients With Rheumatic Disorders. <i>Journal of Bone and Mineral Research</i> , 2015 , 30, 1667-75	6.3	72
230	Intravenous Bisphosphonate Therapy of Young Children With Osteogenesis Imperfecta: Skeletal Findings During Follow Up Throughout the Growing Years. <i>Journal of Bone and Mineral Research</i> , 2015 , 30, 2150-7	6.3	81
229	Mathematical model for bone mineralization. Frontiers in Cell and Developmental Biology, 2015, 3, 51	5.7	12
228	Recessive osteogenesis imperfecta caused by missense mutations in SPARC. <i>American Journal of Human Genetics</i> , 2015 , 96, 979-85	11	81
227	Homozygosity for frameshift mutations in XYLT2 result in a spondylo-ocular syndrome with bone fragility, cataracts, and hearing defects. <i>American Journal of Human Genetics</i> , 2015 , 96, 971-8	11	50
226	The functional muscle-bone unit in patients with osteogenesis imperfecta type I. <i>Bone</i> , 2015 , 79, 52-7	4.7	34
225	Cole-Carpenter syndrome is caused by a heterozygous missense mutation in P4HB. <i>American Journal of Human Genetics</i> , 2015 , 96, 425-31	11	65
224	Unique micro- and nano-scale mineralization pattern of human osteogenesis imperfecta type VI bone. <i>Bone</i> , 2015 , 73, 233-41	4.7	37

223	Physical activity in youth with osteogenesis imperfecta type I. <i>Journal of Musculoskeletal Neuronal Interactions</i> , 2015 , 15, 171-6	1.3	23
222	Effect of High-Dose Vitamin D Supplementation on Bone Density in Youth with Osteogenesis Imperfecta: A Randomized Controlled Trial. <i>FASEB Journal</i> , 2015 , 29, 758.9	0.9	
221	Vitamin D Status and Functional Health Outcomes: A Randomized Vitamin D Dose-Response Trial in 2-8 y Olds. <i>FASEB Journal</i> , 2015 , 29, 734.5	0.9	
220	Milk Product Intake May Not Associate With Body Fat Percentage or Bone Density in Young Children. <i>FASEB Journal</i> , 2015 , 29, 734.9	0.9	
219	Skeletal findings in the first 12 months following initiation of glucocorticoid therapy for pediatric nephrotic syndrome. <i>Osteoporosis International</i> , 2014 , 25, 627-37	5.3	37
218	Muscle anatomy and dynamic muscle function in osteogenesis imperfecta type I. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014 , 99, E356-62	5.6	40
217	Topological mapping of BRIL reveals a type II orientation and effects of osteogenesis imperfecta mutations on its cellular destination. <i>Journal of Bone and Mineral Research</i> , 2014 , 29, 2004-16	6.3	23
216	Pediatric reference Raman data for material characteristics of iliac trabecular bone. <i>Bone</i> , 2014 , 69, 89	-9 7 .7	36
215	Circulating sclerostin in children and young adults with heritable bone disorders. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014 , 99, E920-5	5.6	17
214	Effect of sclerostin antibody treatment in a mouse model of severe osteogenesis imperfecta. <i>Bone</i> , 2014 , 66, 182-8	4.7	73
213	Skeletal characteristics associated with homozygous and heterozygous WNT1 mutations. <i>Bone</i> , 2014 , 67, 63-70	4.7	34
212	Targeted sequencing of a pediatric metabolic bone gene panel using a desktop semiconductor next-generation sequencer. <i>Calcified Tissue International</i> , 2014 , 95, 323-31	3.9	21
211	Shaping and managing the course of a childß disease: parental experiences with osteogenesis imperfecta. <i>Disability and Health Journal</i> , 2014 , 7, 343-9	4.2	18
210	Changes in Bone Density during Development 2014 , 231-235		
209	From pediatric to adult care: strategic evaluation of a transition program for patients with osteogenesis imperfecta. <i>BMC Health Services Research</i> , 2014 , 14, 489	2.9	18
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177 176 175	Low bone mass and high material bone density in two patients with Loeys-Dietz syndrome caused by transforming growth factor beta receptor 2 mutations. <i>Journal of Bone and Mineral Research</i> , 2012 , 27, 713-8 A co-occurrence of osteogenesis imperfecta type VI and cystinosis. <i>American Journal of Medical Genetics, Part A</i> , 2012 , 158A, 1422-6 High incidence of vertebral fractures in children with acute lymphoblastic leukemia 12 months after the initiation of therapy. <i>Journal of Clinical Oncology</i> , 2012 , 30, 2760-7 Lack of circulating pigment epithelium-derived factor is a marker of osteogenesis imperfecta type	2.5	18 15 98
177 176 175	Low bone mass and high material bone density in two patients with Loeys-Dietz syndrome caused by transforming growth factor beta receptor 2 mutations. <i>Journal of Bone and Mineral Research</i> , 2012 , 27, 713-8 A co-occurrence of osteogenesis imperfecta type VI and cystinosis. <i>American Journal of Medical Genetics, Part A</i> , 2012 , 158A, 1422-6 High incidence of vertebral fractures in children with acute lymphoblastic leukemia 12 months after the initiation of therapy. <i>Journal of Clinical Oncology</i> , 2012 , 30, 2760-7 Lack of circulating pigment epithelium-derived factor is a marker of osteogenesis imperfecta type VI. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012 , 97, E1550-6 Serum 24,25-dihydroxyvitamin D concentrations in osteogenesis imperfecta: relationship to bone	2.5 2.2 5.6	18 15 98 53
177 176 175 174	Low bone mass and high material bone density in two patients with Loeys-Dietz syndrome caused by transforming growth factor beta receptor 2 mutations. <i>Journal of Bone and Mineral Research</i> , 2012 , 27, 713-8 A co-occurrence of osteogenesis imperfecta type VI and cystinosis. <i>American Journal of Medical Genetics, Part A</i> , 2012 , 158A, 1422-6 High incidence of vertebral fractures in children with acute lymphoblastic leukemia 12 months after the initiation of therapy. <i>Journal of Clinical Oncology</i> , 2012 , 30, 2760-7 Lack of circulating pigment epithelium-derived factor is a marker of osteogenesis imperfecta type VI. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012 , 97, E1550-6 Serum 24,25-dihydroxyvitamin D concentrations in osteogenesis imperfecta: relationship to bone parameters. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012 , 97, 1243-9 Muscle-bone characteristics in children with Prader-Willi syndrome. <i>Journal of Clinical Endocrinology</i>	2.5 2.2 5.6	18 15 98 53

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141	Osteogenesis imperfecta type III with intracranial hemorrhage and brachydactyly associated with mutations in exon 49 of COL1A2. <i>American Journal of Medical Genetics, Part A</i> , 2009 , 149A, 461-5 Intravenous pamidronate in osteogenesis imperfecta type VII. <i>Calcified Tissue International</i> , 2009 ,		17
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141 140 139	Osteogenesis imperfecta type III with intracranial hemorrhage and brachydactyly associated with mutations in exon 49 of COL1A2. <i>American Journal of Medical Genetics, Part A</i> , 2009 , 149A, 461-5 Intravenous pamidronate in osteogenesis imperfecta type VII. <i>Calcified Tissue International</i> , 2009 , 84, 203-9 Vibration therapy. <i>Developmental Medicine and Child Neurology</i> , 2009 , 51 Suppl 4, 166-8 Large osteoclasts in pediatric osteogenesis imperfecta patients receiving intravenous pamidronate. <i>Journal of Bone and Mineral Research</i> , 2009 , 24, 669-74 Advanced vertebral fracture among newly diagnosed children with acute lymphoblastic leukemia: results of the Canadian Steroid-Associated Osteoporosis in the Pediatric Population (STOPP)	3.9 3.3 6.3	17 12 62 22
141 140 139 138	Osteogenesis imperfecta type III with intracranial hemorrhage and brachydactyly associated with mutations in exon 49 of COL1A2. <i>American Journal of Medical Genetics, Part A</i> , 2009 , 149A, 461-5 Intravenous pamidronate in osteogenesis imperfecta type VII. <i>Calcified Tissue International</i> , 2009 , 84, 203-9 Vibration therapy. <i>Developmental Medicine and Child Neurology</i> , 2009 , 51 Suppl 4, 166-8 Large osteoclasts in pediatric osteogenesis imperfecta patients receiving intravenous pamidronate. <i>Journal of Bone and Mineral Research</i> , 2009 , 24, 669-74 Advanced vertebral fracture among newly diagnosed children with acute lymphoblastic leukemia: results of the Canadian Steroid-Associated Osteoporosis in the Pediatric Population (STOPP) research program. <i>Journal of Bone and Mineral Research</i> , 2009 , 24, 1326-34 Risedronate in the treatment of mild pediatric osteogenesis imperfecta: a randomized	3.9 3.3 6.3	17 12 62 22 151

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