

Laura L Tosi

List of Publications by Year in descending order

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Version: 2024-02-01

32
papers

2,185
citations

430874

18
h-index

414414

32
g-index

38
all docs

38
docs citations

38
times ranked

3448
citing authors

#	ARTICLE	IF	CITATIONS
1	Best Practice Guidelines for Assessment and Management of Osteoporosis in Adult Patients Undergoing Elective Spinal Reconstruction. <i>Spine</i> , 2022, 47, 128-135.	2.0	22
2	Global variations in barriers to exercise for individuals with intellectual and developmental disability. <i>The Lancet Global Health</i> , 2022, 10, S15.	6.3	0
3	Musculoskeletal abnormalities in a large international cohort of boys with 49, <scpx>XXXXY</scpx>. <i>American Journal of Medical Genetics, Part A</i> , 2021, 185, 3531-3540.	1.2	3
4	<scpx>Health-related</scpx> quality of life in adults with osteogenesis imperfecta. <i>Clinical Genetics</i> , 2021, 99, 772-779.	2.0	4
5	A standard set of outcome measures for the comprehensive assessment of osteogenesis imperfecta. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 140.	2.7	18
6	Somatic PIK3R1 variation as a cause of vascular malformations and overgrowth. <i>Genetics in Medicine</i> , 2021, 23, 1882-1888.	2.4	26
7	Secondary Fracture Prevention: Consensus Clinical Recommendations from a Multistakeholder Coalition. <i>Journal of Bone and Mineral Research</i> , 2020, 35, 36-52.	2.8	146
8	Fourteen-year follow-up of a child with acroscyphodysplasia with emphasis on the need for multidisciplinary management: a case report. <i>BMC Medical Genetics</i> , 2020, 21, 189.	2.1	2
9	Secondary Fracture Prevention: Consensus Clinical Recommendations from a Multistakeholder Coalition. <i>Journal of Orthopaedic Trauma</i> , 2020, 34, e125-e141.	1.4	10
10	Malocclusion traits and oral health-related quality of life in children with osteogenesis imperfecta. <i>Journal of the American Dental Association</i> , 2020, 151, 480-490.e2.	1.5	9
11	The Rare Bone Disease TeleECHO Program: Leveraging Telehealth to Improve Rare Bone Disease Care. <i>Current Osteoporosis Reports</i> , 2020, 18, 344-349.	3.6	16
12	Distinct interferon signatures and cytokine patterns define additional systemic autoinflammatory diseases. <i>Journal of Clinical Investigation</i> , 2020, 130, 1669-1682.	8.2	142
13	Assessing disease experience across the life span for individuals with osteogenesis imperfecta: challenges and opportunities for patient-reported outcomes (PROs) measurement: a pilot study. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 23.	2.7	19
14	Incorporating the patient perspective in the study of rare bone disease: insights from the osteogenesis imperfecta community. <i>Osteoporosis International</i> , 2019, 30, 507-511.	3.1	16
15	Genetic characterization of physical activity behaviours in university students enrolled in kinesiology degree programs. <i>Applied Physiology, Nutrition and Metabolism</i> , 2017, 42, 278-284.	1.9	5
16	Own the Bone, a System-Based Intervention, Improves Osteoporosis Care After Fragility Fractures. <i>Journal of Bone and Joint Surgery - Series A</i> , 2016, 98, e109.	3.0	69
17	Initial report of the osteogenesis imperfecta adult natural history initiative. <i>Orphanet Journal of Rare Diseases</i> , 2015, 10, 146.	2.7	46
18	Impaired osteoblast and osteoclast function characterize the osteoporosis of Snyder - Robinson syndrome. <i>Orphanet Journal of Rare Diseases</i> , 2015, 10, 27.	2.7	27

#	ARTICLE	IF	CITATIONS
19	Mechanistic and therapeutic insights gained from studying rare skeletal diseases. <i>Bone</i> , 2015, 76, 67-75.	2.9	22
20	Editorial Comment: Symposium: Sex Differences in Musculoskeletal Disease and Science. <i>Clinical Orthopaedics and Related Research</i> , 2015, 473, 2474-2478.	1.5	5
21	Declining Rates of Osteoporosis Management Following Fragility Fractures in the U.S., 2000 Through 2009. <i>Journal of Bone and Joint Surgery - Series A</i> , 2014, 96, e52.	3.0	96
22	Bisphosphonate Treatment for Children With Disabling Conditions. <i>PM and R</i> , 2014, 6, 427-436.	1.6	44
23	Clinical delineation and natural history of the <i>PIK3CA</i> -related overgrowth spectrum. <i>American Journal of Medical Genetics, Part A</i> , 2014, 164, 1713-1733.	1.2	249
24	A Mosaic Activating Mutation in <i>AKT1</i> Associated with the Proteus Syndrome. <i>New England Journal of Medicine</i> , 2011, 365, 611-619.	27.0	800
25	Bone Quality: Educational Tools for Patients, Physicians, and Educators. <i>Clinical Orthopaedics and Related Research</i> , 2011, 469, 2248-2259.	1.5	6
26	Breakout Session: Sex/Gender and Racial/Ethnic Disparities in the Care of Osteoporosis and Fragility Fractures. <i>Clinical Orthopaedics and Related Research</i> , 2011, 469, 1936-1940.	1.5	8
27	Assessment and management of the orthopedic and other complications of Proteus syndrome. <i>Journal of Children's Orthopaedics</i> , 2011, 5, 319-327.	1.1	33
28	The American Orthopaedic Association's "Own the Bone" Initiative to Prevent Secondary Fractures. <i>Journal of Bone and Joint Surgery - Series A</i> , 2008, 90, 163-173.	3.0	124
29	Does Sex Matter in Musculoskeletal Health? A Workshop Report. <i>Orthopedic Clinics of North America</i> , 2006, 37, 523-529.	1.2	9
30	Does Sex Matter in Musculoskeletal Health? The Influence of Sex and Gender on Musculoskeletal Health. <i>Journal of Bone and Joint Surgery - Series A</i> , 2005, 87, 1631.	3.0	61
31	An AOA Critical Issue. <i>Journal of Bone and Joint Surgery - Series A</i> , 2005, 87, 2812-2821.	3.0	17
32	Three novel type I collagen mutations in osteogenesis imperfecta type IV probands are associated with discrepancies between electrophoretic migration of osteoblast and fibroblast collagen. <i>Human Mutation</i> , 1998, 11, 395-403.	2.5	37