Mauro Celli

List of Publications by Year
in descending order

46	1,033	623188	433756
papers	citations	14 h-index	31 g-index

47	47	47	1312
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Vertebral fracture assessment (VFA) for monitoring vertebral reshaping in children and adolescents with osteogenesis imperfecta treated with intravenous neridronate. Bone, 2021, 143, 115608.	1.4	5
2	Challenges in surgical orthopaedic treatment in a rare case of pycnodysostosis: Sometimes you win, sometimes you learn. Journal of Orthopaedic Science, 2021, , .	0.5	0
3	The Characteristics of Adjacent Anatomy of Mandibular Third Molar Germs: A CBCT Pilot Study in Patients with Osteogenesis Imperfecta. Healthcare (Switzerland), 2020, 8, 372.	1.0	4
4	New 3D Cone Beam CT Imaging Parameters to Assist the Dentist in Treating Patients with Osteogenesis Imperfecta. Healthcare (Switzerland), 2020, 8, 546.	1.0	3
5	Ultrastructure study of skin fibroblasts in patients with Ehlers-Danlos Syndrome (EDS): preliminary results. Clinica Terapeutica, 2020, 171, e431-e436.	0.2	1
6	Genotypeâ€“phenotype correlation study in 364 osteogenesis imperfecta Italian patients. European Journal of Human Genetics, 2019, 27, 1090-1100.	1.4	52
7	Elastic intramedullary nailing of the femur fracture in patients affected by osteogenesis imperfecta type 3: Indications, limits and pitfalls. Injury, 2019, 50, S52-S56.	0.7	13
8	Treatment of tibial deformities with the Fassierâ€“Duval telescopic nail and minimally invasive percutaneous osteotomies in patients with osteogenesis imperfecta type III. Journal of Pediatric Orthopaedics Part B, 2019, 28, 179-185.	0.3	11
9	Effectiveness of oral propranolol in a patient with neurofibromatosis type 1 and infantile hemangiomas. Italian Journal of Dermatology and Venereology, 2019, , .	0.1	1
10	Intraoperative bleeding in patients with osteogenesis imperfecta type III treated by Fassierâ€“Duval femoral rodding: analysis of risk factors. Journal of Pediatric Orthopaedics Part B, 2018, 27, 338-343.	0.3	9
11	Radial head dislocation and malalignment in osteogenesis imperfecta type V: case report, pitfalls in the treatment, and review of the literature. Journal of Pediatric Orthopaedics Part B, 2018, 27, 375-378.	0.3	3
12	Evaluation of temporomandibular disorders and comorbidities in patients with Ehler-â€“Danlos: Clinical and digital findings. Journal of International Society of Preventive and Community Dentistry, 2018, 8, 333.	0.4	8
13	Association between spondylolisthesis and L5 fracture in patients with Osteogenesis Imperfecta. European Spine Journal, 2017, 26, 3106-3111.	1.0	2
14	Serum creatine kinase isoenzymes in children with osteogenesis imperfecta. Osteoporosis International, 2017, 28, 339-346.	1.3	6
15	BISPHOSPHONATES THERAPY IN CHILDREN WITH OSTEOGENESIS IMPERFECTA: CLINICAL EXPERIENCE IN ORAL SURGERY. ORAL and Implantology, 2017, 10, 311.	0.3	7
16	Clinical and biochemical response to neridronate treatment in a patient with osteoporosis-pseudoglioma syndrome (OPPG). Osteoporosis International, 2017, 28, 3277-3280.	1.3	5
17	Creatine Kinase as Biomarker in Osteogenesis Imperfecta. Biomarkers in Disease, 2017, , 177-199.	0.0	0
18	Osteogenesis imperfecta and clubfootâ€”a rare combination. Medicine (United States), 2016, 95, e4505.	0.4	3

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19	Osteochondritis dissecans of the lateral femoral condyle in a patient affected by osteogenesis imperfecta. Journal of Pediatric Orthopaedics Part B, 2015, 24, 521-525.	0.3	13
20	Phenotypic variability in developmental coordination disorder: Clustering of generalized joint hypermobility with attention deficit/hyperactivity disorder, atypical swallowing and narrative difficulties. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2015, 169, 117-122.	0.7	16
21	Osteogenesis imperfecta and rapid maxillary expansion: Report of 3 patients. American Journal of Orthodontics and Dentofacial Orthopedics, 2015, 148, 130-137.	0.8	15
22	Reliability of Vertebral Fractures Assessment (VFA) in Children with Osteogenesis Imperfecta. Calcified Tissue International, 2015, 96, 307-312.	1.5	37
23	Serum brain-type creatine kinase increases in children with osteogenesis imperfecta during neridronate treatment. Pediatric Research, 2014, 75, 626-630.	1.1	8
24	Management of pain and fatigue in the joint hypermobility syndrome (a.k.a. Ehlersâ€“Danlos syndrome,) Tj ETQq0 0 0 rgBT /Overlock 10 Medical Genetics, Part A, 2012, 158A, 2055-2070.	0.7	124
25	Vitamin D deficiency rickets in five â€œatâ€riskâ€•children. Pediatrics International, 2012, 54, 152-155.	0.2	3
26	Tyrosinemia type I: long-term outcome in a patient treated with doses of NTBC lower than recommended. European Journal of Pediatrics, 2011, 170, 819-819.	1.3	6
27	Osteogenesis imperfecta: the audiological phenotype lacks correlation with the genotype. Orphanet Journal of Rare Diseases, 2011, 6, 88.	1.2	48
28	Taurine deficiency in thalassemia major-induced osteoporosis treated with neridronate. Biomedicine and Pharmacotherapy, 2010, 64, 271-274.	2.5	7
29	Increased nitric oxide release by neutrophils of a patient with tyrosinemia type III. Biomedicine and Pharmacotherapy, 2009, 63, 359-361.	2.5	8
30	Impairment of diastolic function in adult patients affected by osteogenesis imperfecta clinically asymptomatic for cardiac disease: Casuality or causality?. International Journal of Cardiology, 2009, 131, 200-203.	0.8	33
31	High Levels of Serum Prostaglandin E2 in Children with Osteogenesis Imperfecta Are Reduced by Neridronate Treatment. Pediatric Research, 2008, 63, 203-206.	1.1	14
32	Reduction of plasma taurine level in children affected by Osteogenesis Imperfecta during bisphosphonate therapy. Biomedicine and Pharmacotherapy, 2007, 61, 235-240.	2.5	7
33	Absence of severe recurrent infections in glycogen storage disease type Ib with neutropenia and neutrophil dysfunction. Journal of Inherited Metabolic Disease, 2007, 30, 105-105.	1.7	1
34	Neutrophil Glutamine Deficiency in Relation to Genotype in Children with Cystic Fibrosis. Pediatric Research, 2006, 59, 13-16.	1.1	18
35	Fatty acid profile of oesophageal mucosa in children with gastro-oesophageal reflux disease. Digestive and Liver Disease, 2003, 35, 694-700.	0.4	1
36	Increased taurine content in esophageal mucosa of children affected by gastroesophageal reflux. Digestive Diseases and Sciences, 2001, 46, 808-814.	1.1	3

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37	Low-Dosage Immunoglobulins for an Infant With Hypogammaglobulinemia, Maple Syrup Urine Disease, and Parvovirus B19-associated Aplastic Crisis. The American Journal of Pediatric Hematology/oncology, 2000, 22, 485-487.	1.3	3
38	Determination of urinary orotic acid and uracil by capillary zone electrophoresis. Biomedical Applications, 1999, 734, 175-178.	1.7	24
39	Effect of d-ribose on purine synthesis and neurological symptoms in a patient with adenylosuccinase deficiency. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 1999, 1453, 135-140.	1.8	34
40	Usefulness of cyanide-nitroprusside test in detecting incomplete recessive heterozygotes for cystinuria: a standardized dilution procedure. Urological Research, 1998, 26, 401-405.	1.5	31
41	Erythrocyte and Plasma Levels of Glutamate and Aspartate in Children Affected by Migraine. Cephalalgia, 1997, 17, 652-657.	1.8	23
42	Abnormal intestinal permeability in children with autism. Acta Paediatrica, International Journal of Paediatrics, 1996, 85, 1076-1079.	0.7	330
43	Immunological abnormalities in a patient with tyrosinaemia type III. Journal of Inherited Metabolic Disease, 1995, 18, 355-356.	1.7	3
44	Low serum tryptophan to large neutral amino acids ratio in idiopathic infantile autism. Biomedicine and Pharmacotherapy, 1995, 49, 288-292.	2.5	48
45	Late onset of cystinuria in a case of gyrate atrophy. Journal of Inherited Metabolic Disease, 1993, 16, 904-905.	1.7	0
46	Child with manifestations of dermotrichic syndrome and ichthyosis follicularis-alopecia-photophobia (IFAP) syndrome. American Journal of Medical Genetics Part A, 1992, 44, 233-236.	2.4	42