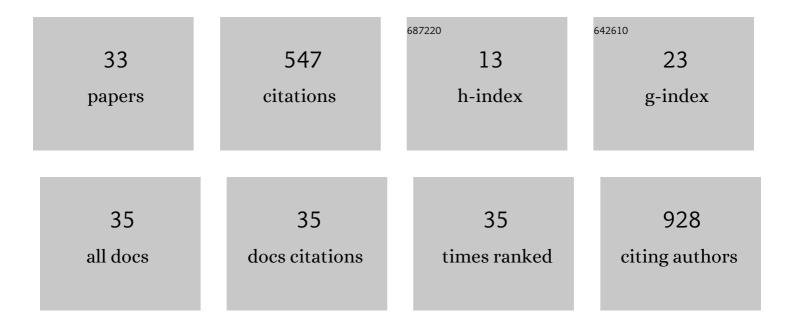
Marta Bertamino

List of Publications by Year in descending order

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MADTA REDTAMINO

#	Article	IF	CITATIONS
1	Executive functions and psychosocial impairment in children following arterial ischemic stroke. Child Neuropsychology, 2023, 29, 276-298.	0.8	1
2	An Atypical Case of Aphasia: Transitory Ischemic Attack in a 13-Year-Old Patient with Asymptomatic SARS-CoV-2 Infection. Children, 2022, 9, 983.	0.6	0
3	Genotype-Phenotype Correlation and Functional Insights for Two Monoallelic TREX1 Missense Variants Affecting the Catalytic Core. Genes, 2022, 13, 1179.	1.0	2
4	Improved trunk and neck control after selective dorsal rhizotomy in children with spastic cerebral palsy. Child's Nervous System, 2021, 37, 351-352.	0.6	1
5	Spontaneous movements in the newborns: a tool of quantitative video analysis of preterm babies. Computer Methods and Programs in Biomedicine, 2021, 199, 105838.	2.6	13
6	An atypical case of post-varicella stroke in a child presenting with hemichorea followed by late-onset inflammatory focal cerebral arteriopathy. Quantitative Imaging in Medicine and Surgery, 2021, 11, 463-471.	1.1	6
7	Communication-vulnerable pediatric patients following posterior fossa tumor surgery: the importance of augmentative and alternative communication. Child's Nervous System, 2021, 37, 2437-2438.	0.6	1
8	Expanding the clinical and neuroimaging features of post-varicella arteriopathy of childhood. Journal of Neurology, 2021, 268, 4846-4865.	1.8	6
9	Combined medical therapy and neurosurgical revascularization preventing stroke in post-varicella angiopathy: Case report and review of literature. Brain and Development, 2021, 43, 1051-1056.	0.6	1
10	Bivalirudin anticoagulation to overcome heparin resistance in a neonate with cerebral sinovenus thrombosis. Blood Coagulation and Fibrinolysis, 2020, 31, 97-100.	0.5	5
11	Targeted re-sequencing in pediatric and perinatal stroke. European Journal of Medical Genetics, 2020, 63, 104030.	0.7	9
12	Perinatal Arterial Ischemic Stroke in Fetal Vascular Malperfusion: A Case Series and Literature Review. American Journal of Neuroradiology, 2020, 41, 2377-2383.	1.2	6
13	Letter to the editor regarding: "High prevalence of pro-thrombotic conditions in adult patients with moyamoya disease and moyamoya syndrome: a single center study― Acta Neurochirurgica, 2020, 162, 3139-3140.	0.9	1
14	Letter to the Editor Regarding "Large Craniotomy Increases the Risk of Minor Perioperative Complications in Revascularization Surgery for Moyamoya Disease― World Neurosurgery, 2020, 143, 581-583.	0.7	3
15	Spatial coefficient of variation applied to arterial spin labeling MRI may contribute to predict surgical revascularization outcomes in pediatric moyamoya vasculopathy. Neuroradiology, 2020, 62, 1003-1015.	1.1	11
16	Schimke immuno-osseous dysplasia, two new cases with peculiar EEG pattern. Brain and Development, 2020, 42, 408-413.	0.6	2
17	Impact on rehabilitation programs during COVID-19 containment for children with pediatric and perinatal stroke. European Journal of Physical and Rehabilitation Medicine, 2020, 56, 692-694.	1.1	15
18	A prediction rule for lack of achievement of inactive disease with methotrexate as the sole disease-modifying antirheumatic therapy in juvenile idiopathic arthritis. Pediatric Rheumatology, 2019, 17, 50.	0.9	5

MARTA BERTAMINO

#	Article	IF	CITATIONS
19	Consensus statements on vaccination in patients with haemophilia—Results from the Italian haemophilia and vaccinations (HEVA) project. Haemophilia, 2019, 25, 656-667.	1.0	16
20	ABCC6 mutations and early onset stroke: Two cases of a typical Pseudoxanthoma Elasticum. European Journal of Paediatric Neurology, 2018, 22, 725-728.	0.7	15
21	Novel spondyloepimetaphyseal dysplasia due to <i>UFSP2</i> gene mutation. Clinical Genetics, 2018, 93, 671-674.	1.0	26
22	Long term substrate reduction therapy with ezetimibe alone or associated with statins in three adult patients with lysosomal acid lipase deficiency. Orphanet Journal of Rare Diseases, 2018, 13, 24.	1.2	12
23	Widening the Heterogeneity of Leigh Syndrome: Clinical, Biochemical, and Neuroradiologic Features in a Patient Harboring a NDUFA10 Mutation. JIMD Reports, 2017, 37, 37-43.	0.7	13
24	A new form of IRIDA due to combined heterozygous mutations of TMPRSS6 and ACVR1A encoding the BMP receptor ALK2. Blood, 2017, 129, 3392-3395.	0.6	18
25	International physician survey on management of FOP: a modified Delphi study. Orphanet Journal of Rare Diseases, 2017, 12, 110.	1.2	15
26	Hemophilia Care in the Pediatric Age. Journal of Clinical Medicine, 2017, 6, 54.	1.0	39
27	Novel asymptomatic CNS findings in patients withACVR1/ALK2mutations causing fibrodysplasia ossificans progressiva. Journal of Medical Genetics, 2016, 53, 859-864.	1.5	12
28	New insights into central nervous system involvement in FOP: Case report and review of the literature. American Journal of Medical Genetics, Part A, 2015, 167, 2817-2821.	0.7	12
29	A Practical Approach to the Use of Low Molecular Weight Heparins in VTE Treatment and Prophylaxis in Children and Newborns. Pediatric Hematology and Oncology, 2015, 32, 1-10.	0.3	26
30	Outcome and predicting factors of single and multiple intra-articular corticosteroid injections in children with juvenile idiopathic arthritis. Rheumatology, 2011, 50, 1627-1634.	0.9	59
31	Evaluation of 21-Numbered Circle and 10-Centimeter Horizontal Line Visual Analog Scales for Physician and Parent Subjective Ratings in Juvenile Idiopathic Arthritis. Journal of Rheumatology, 2010, 37, 1534-1541.	1.0	119
32	Development and Initial Validation of a Radiographic Scoring System for the Hip in Juvenile Idiopathic Arthritis. Journal of Rheumatology, 2010, 37, 432-439.	1.0	35
33	A new short and simple health-related quality of life measurement for paediatric rheumatic diseases: initial validation in juvenile idiopathic arthritis. Rheumatology, 2010, 49, 1272-1280.	0.9	39