Giulia Amico

List of Publications by Year
in descending order

11 papers	164 citations	1684188 5 h-index	1474206 9 g-index
11 all docs	11 docs citations	11 times ranked	353 citing authors

#	Article	IF	Citations
1	Genotype-Phenotype Correlation and Functional Insights for Two Monoallelic TREX1 Missense Variants Affecting the Catalytic Core. Genes, 2022, 13, 1179.	2.4	2
2	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.	2.0	6
3	Expanding the clinical and neuroimaging features of post-varicella arteriopathy of childhood. Journal of Neurology, 2021, 268, 4846-4865.	3.6	6
4	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.	1.1	1
5	The Genetic Landscape of Patent Foramen Ovale: A Systematic Review. Genes, 2021, 12, 1953.	2.4	5
6	Prenatal Diagnosis of an Uncommon 48, XX ,+18+21 Karyotype in a Fetus With Malformations Typical of Both Trisomies. Journal of Ultrasound in Medicine, 2020, 39, 2277-2279.	1.7	0
7	Unravelling the Regions of Mutant F508del-CFTR More Susceptible to the Action of Four Cystic Fibrosis Correctors. International Journal of Molecular Sciences, 2019, 20, 5463.	4.1	15
8	Cryopyrin-associated Periodic Syndromes in Italian Patients: Evaluation of the Rate of Somatic NLRP3 Mosaicism and Phenotypic Characterization. Journal of Rheumatology, 2017, 44, 1667-1673.	2.0	28
9	In vitro recapitulation of the site-specific editing (to wild-type) of mutant IDS mRNA transcripts, and the characterization of IDS protein translated from the edited mRNAs. Human Mutation, 2017, 38, 849-862.	2.5	0
10	Mutation Update ofARSAandPSAPGenes Causing Metachromatic Leukodystrophy. Human Mutation, 2016, 37, 16-27.	2.5	96
11	MLPA-based approach for initial and simultaneous detection of GBA deletions and recombinant alleles in patients affected by Gaucher Disease. Molecular Genetics and Metabolism, 2016, 119, 329-337.	1.1	5