

Giulia Amico

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6342924/publications.pdf>

Version: 2024-02-01

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	Mutation Update of <i>ARSA</i> and <i>PSAP</i> Genes Causing Metachromatic Leukodystrophy. <i>Human Mutation</i> , 2016, 37, 16-27.	2.5	96
2	Cryopyrin-associated Periodic Syndromes in Italian Patients: Evaluation of the Rate of Somatic NLRP3 Mosaicism and Phenotypic Characterization. <i>Journal of Rheumatology</i> , 2017, 44, 1667-1673.	2.0	28
3	Unravelling the Regions of Mutant F508del-CFTR More Susceptible to the Action of Four Cystic Fibrosis Correctors. <i>International Journal of Molecular Sciences</i> , 2019, 20, 5463.	4.1	15
4	An atypical case of post-varicella stroke in a child presenting with hemichorea followed by late-onset inflammatory focal cerebral arteriopathy. <i>Quantitative Imaging in Medicine and Surgery</i> , 2021, 11, 463-471.	2.0	6
5	Expanding the clinical and neuroimaging features of post-varicella arteriopathy of childhood. <i>Journal of Neurology</i> , 2021, 268, 4846-4865.	3.6	6
6	MLPA-based approach for initial and simultaneous detection of GBA deletions and recombinant alleles in patients affected by Gaucher Disease. <i>Molecular Genetics and Metabolism</i> , 2016, 119, 329-337.	1.1	5
7	The Genetic Landscape of Patent Foramen Ovale: A Systematic Review. <i>Genes</i> , 2021, 12, 1953.	2.4	5
8	Genotype-Phenotype Correlation and Functional Insights for Two Monoallelic TREX1 Missense Variants Affecting the Catalytic Core. <i>Genes</i> , 2022, 13, 1179.	2.4	2
9	Combined medical therapy and neurosurgical revascularization preventing stroke in post-varicella angiopathy: Case report and review of literature. <i>Brain and Development</i> , 2021, 43, 1051-1056.	1.1	1
10	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. <i>Human Mutation</i> , 2017, 38, 849-862.	2.5	0
11	Prenatal Diagnosis of an Uncommon 48, XX,+18+21 Karyotype in a Fetus With Malformations Typical of Both Trisomies. <i>Journal of Ultrasound in Medicine</i> , 2020, 39, 2277-2279.	1.7	0