

Arunabha Ghosh

List of Publications by Year in descending order

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Version: 2024-02-01

12
papers

296
citations

1039880

9
h-index

1199470

12
g-index

12
all docs

12
docs citations

12
times ranked

580
citing authors

#	ARTICLE	IF	CITATIONS
1	Enzyme replacement therapy prior to haematopoietic stem cell transplantation in Mucopolysaccharidosis Type I: 10year combined experience of 2 centres. <i>Molecular Genetics and Metabolism</i> , 2016, 117, 373-377.	0.5	51
2	Diagnosing childhood-onset inborn errors of metabolism by next-generation sequencing. <i>Archives of Disease in Childhood</i> , 2017, 102, 1019-1029.	1.0	43
3	Metabolism of Non-Enzymatically Derived Oxysterols: Clues from sterol metabolic disorders. <i>Free Radical Biology and Medicine</i> , 2019, 144, 124-133.	1.3	39
4	Recognition, assessment and management of hypoglycaemia in childhood. <i>Archives of Disease in Childhood</i> , 2016, 101, 575-580.	1.0	36
5	Haematopoietic stem cell gene therapy with γ -irradiation rescues cognitive loss in mucopolysaccharidosis type IIIA. <i>EMBO Molecular Medicine</i> , 2020, 12, e11185.	3.3	31
6	Recommendations on clinical trial design for treatment of Mucopolysaccharidosis Type III. <i>Orphanet Journal of Rare Diseases</i> , 2017, 12, 117.	1.2	27
7	Identification of unusual oxysterols and bile acids with 7-oxo or 3 β ,5 α ,6 β -trihydroxy functions in human plasma by charge-tagging mass spectrometry with multistage fragmentation. <i>Journal of Lipid Research</i> , 2018, 59, 1058-1070.	2.0	21
8	<i>IDUA</i> mutational profile and genotype-phenotype relationships in UK patients with Mucopolysaccharidosis Type I. <i>Human Mutation</i> , 2017, 38, 1555-1568.	1.1	16
9	Phenotypic Heterogeneity in a Congenital Disorder of Glycosylation Caused by Mutations in <i>STT3A</i> . <i>Journal of Child Neurology</i> , 2017, 32, 560-565.	0.7	10
10	Strategies for the Induction of Immune Tolerance to Enzyme Replacement Therapy in Mucopolysaccharidosis Type I. <i>Molecular Therapy - Methods and Clinical Development</i> , 2019, 13, 321-333.	1.8	9
11	Coenzyme Q10 and Pyridoxal Phosphate Deficiency Is a Common Feature in Mucopolysaccharidosis Type III. <i>JIMD Reports</i> , 2015, 25, 1-7.	0.7	8
12	The effect of haemopoietic stem cell transplantation on the ocular phenotype in mucopolysaccharidosis type I (Hurler). <i>Acta Ophthalmologica</i> , 2018, 96, 494-498.	0.6	5