

Arunabha Ghosh

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

12
papers

187
citations

8
h-index

12
g-index

12
ext. papers

249
ext. citations

4.4
avg, IF

2.46
L-index

#	Paper	IF	Citations
12	Haematopoietic stem cell gene therapy with IL-1Ra rescues cognitive loss in mucopolysaccharidosis IIIA. <i>EMBO Molecular Medicine</i> , 2020 , 12, e11185	12	11
11	Metabolism of Non-Enzymatically Derived Oxysterols: Clues from sterol metabolic disorders. <i>Free Radical Biology and Medicine</i> , 2019 , 144, 124-133	7.8	19
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	6.4	5
9	Identification of unusual oxysterols and bile acids with 7-oxo or 3 β ,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	6.3	20
8	The effect of haematopoietic stem cell transplantation on the ocular phenotype in mucopolysaccharidosis type I (Hurler). <i>Acta Ophthalmologica</i> , 2018 , 96, 494-498	3.7	4
7	Phenotypic Heterogeneity in a Congenital Disorder of Glycosylation Caused by Mutations in STT3A. <i>Journal of Child Neurology</i> , 2017 , 32, 560-565	2.5	8
6	Diagnosing childhood-onset inborn errors of metabolism by next-generation sequencing. <i>Archives of Disease in Childhood</i> , 2017 , 102, 1019-1029	2.2	24
5	IDUA mutational profile and genotype-phenotype relationships in UK patients with Mucopolysaccharidosis Type I. <i>Human Mutation</i> , 2017 , 38, 1555-1568	4.7	12
4	Recommendations on clinical trial design for treatment of Mucopolysaccharidosis Type III. <i>Orphanet Journal of Rare Diseases</i> , 2017 , 12, 117	4.2	16
3	Coenzyme Q and Pyridoxal Phosphate Deficiency Is a Common Feature in Mucopolysaccharidosis Type III. <i>JIMD Reports</i> , 2016 , 25, 1-7	1.9	4
2	Enzyme replacement therapy prior to haematopoietic stem cell transplantation in Mucopolysaccharidosis Type I: 10 year combined experience of 2 centres. <i>Molecular Genetics and Metabolism</i> , 2016 , 117, 373-7	3.7	42
1	Recognition, assessment and management of hypoglycaemia in childhood. <i>Archives of Disease in Childhood</i> , 2016 , 101, 575-580	2.2	22