Arunabha Ghosh

List of Publications by Citations

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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	Enzyme replacement therapy prior to haematopoietic stem cell transplantation in Mucopolysaccharidosis Type I: 10 year combined experience of 2 centres. *Molecular Genetics and Metabolism*, **2016**, 117, 373-7	3.7	42
11	Diagnosing childhood-onset inborn errors of metabolism by next-generation sequencing. *Archives of Disease in Childhood*, **2017**, 102, 1019-1029	2.2	24
10	Recognition, assessment and management of hypoglycaemia in childhood. *Archives of Disease in Childhood*, **2016**, 101, 575-580	2.2	22
9	Identification of unusual oxysterols and bile acids with 7-oxo or 3,5,6-trihydroxy functions in human plasma by charge-tagging mass spectrometry with multistage fragmentation. *Journal of Lipid Research*, **2018**, 59, 1058-1070	6.3	20
8	Metabolism of Non-Enzymatically Derived Oxysterols: Clues from sterol metabolic disorders. *Free Radical Biology and Medicine*, **2019**, 144, 124-133	7.8	19
7	Recommendations on clinical trial design for treatment of Mucopolysaccharidosis Type III. *Orphanet Journal of Rare Diseases*, **2017**, 12, 117	4.2	16
6	IDUA mutational profile and genotype-phenotype relationships in UK patients with Mucopolysaccharidosis Type I. *Human Mutation*, **2017**, 38, 1555-1568	4.7	12
5	Haematopoietic stem cell gene therapy with IL-1Ra rescues cognitive loss in mucopolysaccharidosis IIIA. *EMBO Molecular Medicine*, **2020**, 12, e11185	12	11
4	Phenotypic Heterogeneity in a Congenital Disorder of Glycosylation Caused by Mutations in STT3A. *Journal of Child Neurology*, **2017**, 32, 560-565	2.5	8
3	Strategies for the Induction of Immune Tolerance to Enzyme Replacement Therapy in Mucopolysaccharidosis Type I. *Molecular Therapy - Methods and Clinical Development*, **2019**, 13, 321-333	6.4	5
2	Coenzyme Q and Pyridoxal Phosphate Deficiency Is a Common Feature in Mucopolysaccharidosis Type III. *JIMD Reports*, **2016**, 25, 1-7	1.9	4
1	The effect of haemopoietic stem cell transplantation on the ocular phenotype in mucopolysaccharidosis type I (Hurler). *Acta Ophthalmologica*, **2018**, 96, 494-498	3.7	4