Gouri Yogalingam

List of Publications by Year in descending order

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933447 1281871 12 472 10 11 citations g-index h-index papers 13 13 13 647 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Molecular genetics of mucopolysaccharidosis type IIIA and IIIB: Diagnostic, clinical, and biological implications. Human Mutation, 2001, 18, 264-281.	2.5	153
2	Neuraminidase 1 Is a Negative Regulator of Lysosomal Exocytosis. Developmental Cell, 2008, 15, 74-86.	7.0	136
3	Clearance of Heparan Sulfate and Attenuation of CNS Pathology by Intracerebroventricular BMN 250 in Sanfilippo Type B Mice. Molecular Therapy - Methods and Clinical Development, 2017, 6, 43-53.	4.1	34
4	Intracerebroventricular enzyme replacement therapy with \hat{l}^2 -galactosidase reverses brain pathologies due to GM1 gangliosidosis in mice. Journal of Biological Chemistry, 2020, 295, 13532-13555.	3.4	30
5	Mucopolysaccharidosis type IIIB: characterisation and expression of wild-type and mutant recombinant α-N-acetylglucosaminidase and relationship with Sanfilippo phenotype in an attenuated patient. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2000, 1502, 415-425.	3.8	29
6	Mild Feline Mucopolysaccharidosis Type VI. Journal of Biological Chemistry, 1998, 273, 13421-13429.	3.4	18
7	BMN 250, a fusion of lysosomal alpha-N-acetylglucosaminidase with IGF2, exhibits different patterns of cellular uptake into critical cell types of Sanfilippo syndrome B disease pathogenesis. PLoS ONE, 2019, 14, e0207836.	2.5	18
8	Cellular Uptake and Delivery of Myeloperoxidase to Lysosomes Promote Lipofuscin Degradation and Lysosomal Stress in Retinal Cells. Journal of Biological Chemistry, 2017, 292, 4255-4265.	3.4	17
9	Characterization of glycan substrates accumulating in GM1 Gangliosidosis. Molecular Genetics and Metabolism Reports, 2019, 21, 100524.	1.1	15
10	Differential Uptake of NAGLU-IGF2 and Unmodified NAGLU in Cellular Models of Sanfilippo Syndrome Type B. Molecular Therapy - Methods and Clinical Development, 2019, 14, 56-63.	4.1	12
11	Intermittent enzyme replacement therapy with recombinant human \hat{l}^2 -galactosidase prevents neuraminidase 1 deficiency. Journal of Biological Chemistry, 2020, 295, 13556-13569.	3.4	10
12	Aryplase (Biomarin). Current Opinion in Investigational Drugs, 2004, 5, 1111-20.	2.3	0