Alia Ahmed

List of Publications by Year
in descending order

14 papers	321 citations	933447 10 h-index	996975 15 g-index
15 all docs	15 docs citations	15 times ranked	383 citing authors

#	Article	IF	Citations
1	Quantitative brain MRI morphology in severe and attenuated forms of mucopolysaccharidosis type I. Molecular Genetics and Metabolism, 2022, 135, 122-132.	1.1	5
2	Quantifying medical manifestations in hurler syndrome with the infant physical symptom score: Associations with long-term physical and adaptive outcomes. Molecular Genetics and Metabolism, 2022, , .	1.1	2
3	A longitudinal study of neurocognition and behavior in patients with Hurler-Scheie syndrome heterozygous for the L238Q mutation. Molecular Genetics and Metabolism Reports, 2019, 20, 100484.	1.1	3
4	Observing the advanced disease course in mucopolysaccharidosis, type IIIA; a case series. Molecular Genetics and Metabolism, 2018, 123, 123-126.	1.1	18
5	Distinct progression patterns of brain disease in infantile and juvenile gangliosidoses: Volumetric quantitative MRI study. Molecular Genetics and Metabolism, 2018, 123, 97-104.	1.1	35
6	Intrathecal enzyme replacement therapy reverses cognitive decline in mucopolysaccharidosis type I. American Journal of Medical Genetics, Part A, 2017, 173, 780-783.	1.2	26
7	Long-term cognitive and somatic outcomes of enzyme replacement therapy in untransplanted Hurler syndrome. Molecular Genetics and Metabolism Reports, 2017, 13, 64-68.	1.1	18
8	A longitudinal study of emotional adjustment, quality of life and adaptive function in attenuated MPS II. Molecular Genetics and Metabolism Reports, 2016, 7, 32-39.	1.1	32
9	Association of somatic burden of disease with age and neuropsychological measures in attenuated mucopolysaccharidosis types I, II and VI. Molecular Genetics and Metabolism Reports, 2016, 7, 27-31.	1.1	20
10	Quantifying behaviors of children with Sanfilippo syndrome: The Sanfilippo Behavior Rating Scale. Molecular Genetics and Metabolism, 2015, 114, 594-598.	1.1	31
11	Clinical outcomes of Hurler syndrome treated exclusively with enzyme replacement therapy from a young age. Molecular Genetics and Metabolism, 2015, 114, S40.	1.1	2
12	Cognitive, medical, and neuroimaging characteristics of attenuated mucopolysaccharidosis type II. Molecular Genetics and Metabolism, 2015, 114, 170-177.	1.1	43
13	Neurocognition across the spectrum of mucopolysaccharidosis type I: Age, severity, and treatment. Molecular Genetics and Metabolism, 2015, 116, 61-68.	1.1	59
14	Neurocognitive and neuropsychiatric phenotypes associated with the mutation L238Q of the Î±-L-iduronidase gene in Hurlerâ€“Scheie syndrome. Molecular Genetics and Metabolism, 2014, 111, 123-127.	1.1	26