

Kathleen A Delaney

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1659348/publications.pdf>

Version: 2024-02-01

11
papers

408
citations

933447

10
h-index

1281871

11
g-index

11
all docs

11
docs citations

11
times ranked

428
citing authors

#	ARTICLE	IF	CITATIONS
1	Quantitative brain MRI morphology in severe and attenuated forms of mucopolysaccharidosis type I. <i>Molecular Genetics and Metabolism</i> , 2022, 135, 122-132.	1.1	5
2	Cognitive and adaptive measurement endpoints for clinical trials in mucopolysaccharidoses types I, II, and III: A review of the literature. <i>Molecular Genetics and Metabolism</i> , 2017, 121, 57-69.	1.1	20
3	Assessments of neurocognitive and behavioral function in the mucopolysaccharidoses. <i>Molecular Genetics and Metabolism</i> , 2017, 122, 8-16.	1.1	44
4	A Prospective Natural History Study of Mucopolysaccharidosis Type IIIA. <i>Journal of Pediatrics</i> , 2016, 170, 278-287.e4.	1.8	91
5	Neurocognitive clinical outcome assessments for inborn errors of metabolism and other rare conditions. <i>Molecular Genetics and Metabolism</i> , 2016, 118, 65-69.	1.1	28
6	A longitudinal study of emotional adjustment, quality of life and adaptive function in attenuated MPS II. <i>Molecular Genetics and Metabolism Reports</i> , 2016, 7, 32-39.	1.1	32
7	Quantifying behaviors of children with Sanfilippo syndrome: The Sanfilippo Behavior Rating Scale. <i>Molecular Genetics and Metabolism</i> , 2015, 114, 594-598.	1.1	31
8	Cognitive, medical, and neuroimaging characteristics of attenuated mucopolysaccharidosis type II. <i>Molecular Genetics and Metabolism</i> , 2015, 114, 170-177.	1.1	43
9	Neurocognitive and neuropsychiatric phenotypes associated with the mutation L238Q of the Î±-L-iduronidase gene in Hurler-Scheie syndrome. <i>Molecular Genetics and Metabolism</i> , 2014, 111, 123-127.	1.1	26
10	Acquired Autistic Behaviors in Children with Mucopolysaccharidosis Type IIIA. <i>Journal of Pediatrics</i> , 2014, 164, 1147-1151.e1.	1.8	41
11	Methods of Neurodevelopmental Assessment in Children with Neurodegenerative Disease: Sanfilippo Syndrome. <i>JIMD Reports</i> , 2013, 13, 129-137.	1.5	47