# Alia Ahmed

List of Publications by Year
in descending order



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|---|---|---|---|
| 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 | Neurocognition across the spectrum of mucopolysaccharidosis type I: Age, severity, and treatment. Molecular Genetics and Metabolism, 2015, 116, 61-68. | 1.1 | 59 |
| 2 | Cognitive, medical, and neuroimaging characteristics of attenuated mucopolysaccharidosis type II. Molecular Genetics and Metabolism, 2015, 114, 170-177. | 1.1 | 43 |
| 3 | 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 |
| 4 | 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 |
| 5 | Quantifying behaviors of children with Sanfilippo syndrome: The Sanfilippo Behavior Rating Scale. Molecular Genetics and Metabolism, 2015, 114, 594-598. | 1.1 | 31 |
| 6 | 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 |
| 7 | 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 |
| 8 | 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 |
| 9 | Observing the advanced disease course in mucopolysaccharidosis, type IIIA; a case series. Molecular Genetics and Metabolism, 2018, 123, 123-126. | 1.1 | 18 |
| 10 | 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 |
| 11 | Quantitative brain MRI morphology in severe and attenuated forms of mucopolysaccharidosis type I. Molecular Genetics and Metabolism, 2022, 135, 122-132. | 1.1 | 5 |
| 12 | 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 |
| 13 | 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 |
| 14 | 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 |