Shehla Mohammed

List of Publications by Year
in descending order

6	249	1937685 4	1872680 6
papers	citations	h-index	g-index

7	7	7	860
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	RAC1 Missense Mutations in Developmental Disorders with Diverse Phenotypes. American Journal of Human Genetics, 2017, 101, 466-477.	6.2	119
2	De novo, heterozygous, lossâ€ofâ€function mutations in SYNGAP1 cause a syndromic form of intellectual disability. American Journal of Medical Genetics, Part A, 2015, 167, 2231-2237.	1.2	96
3	Recessive MYH7-related myopathy in two families. Neuromuscular Disorders, 2019, 29, 456-467.	0.6	14
4	Childhood-onset generalized epilepsy in Bainbridge-Ropers syndrome. Epilepsy Research, 2018, 140, 166-170.	1.6	12
5	Growth charts in Cockayne syndrome type 1 and type 2. European Journal of Medical Genetics, 2021, 64, 104105.	1.3	4
6	Generation of an iPSC line (CRICKi001-A) from an individual with a germline SMARCA4 missense mutation and autism spectrum disorder. Stem Cell Research, 2021, 53, 102304.	0.7	4