Hisham Megahed

List of Publications by Year
in descending order

8 papers	597 citations	1683934 5 h-index	1719901 7 g-index
9 all docs	9 docs citations	9 times ranked	1849 citing authors

#	Article	IF	Citations
1	Exome Sequencing Links Corticospinal Motor Neuron Disease to Common Neurodegenerative Disorders. Science, 2014, 343, 506-511.	6.0	466
2	Biallelic loss of human CTNNA2, encoding Î±N-catenin, leads to ARP2/3 complex overactivity and disordered cortical neuronal migration. Nature Genetics, 2018, 50, 1093-1101.	9.4	70
3	Utility of whole exome sequencing for the early diagnosis of pediatric-onset cerebellar atrophy associated with developmental delay in an inbred population. Orphanet Journal of Rare Diseases, 2016, 11, 57.	1.2	31
4	Inherited glycosylphosphatidylinositol defects cause the rare Emm-negative blood phenotype and developmental disorders. Blood, 2021, 137, 3660-3669.	0.6	18
5	Clinical and genetic characterization of ten Egyptian patients with Wolfâ€“Hirschhorn syndrome and review of literature. Molecular Genetics & Genomic Medicine, 2021, 9, e1546.	0.6	6
6	Association of GSTP1 p.Ile105Val (rs1695, c.313Aâ€‰>â€‰G) Variant with the Risk of Breast Carcinoma among Egyptian Women. Biochemical Genetics, 2021, 59, 1487-1505.	0.8	5
7	Serum homocysteine, lipid profile and BMI as atherosclerotic risk factors in children with numerical chromosomal aberrations. World Journal of Pediatrics, 2022, 18, 443-448.	0.8	1
8	Clinical Implications of S100A12 and Resolvin D1 Serum Levels, and Related Genes in Children with Familial Mediterranean Fever. Journal of Child Science, 2021, 11, e163-e169.	0.1	0