Hisham Megahed

List of Publications by Year in Descending Order

8 papers	452 citations	4 h-index	9 g-index
9 ext. papers	536 ext. citations	10.3 avg, IF	1.57 L-index

#	Paper	IF	Citations
8	Serum homocysteine, lipid profile and BMI as atherosclerotic risk factors in children with numerical chromosomal aberrations.. *World Journal of Pediatrics*, **2022**, 1	4.6	1
7	Inherited glycosylphosphatidylinositol defects cause the rare Emm-negative blood phenotype and developmental disorders. *Blood*, **2021**, 137, 3660-3669	2.2	7
6	Association of GSTP1 p.Ile105Val (rs1695, c.313A > G) Variant with the Risk of Breast Carcinoma among Egyptian Women. *Biochemical Genetics*, **2021**, 59, 1487-1505	2.4	1
5	Clinical and genetic characterization of ten Egyptian patients with Wolf-Hirschhorn syndrome and review of literature. *Molecular Genetics & Genomic Medicine*, **2021**, 9, e1546	2.3	1
4	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	0.2	
3	Biallelic loss of human CTNNA2, encoding N-catenin, leads to ARP2/3 complex overactivity and disordered cortical neuronal migration. *Nature Genetics*, **2018**, 50, 1093-1101	36.3	48
2	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	4.2	20
1	Exome sequencing links corticospinal motor neuron disease to common neurodegenerative disorders. *Science*, **2014**, 343, 506-511	33.3	374