

Hisham Megahed

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

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	Exome sequencing links corticospinal motor neuron disease to common neurodegenerative disorders. <i>Science</i> , 2014 , 343, 506-511	33.3	374
7	Biallelic loss of human CTNNA2, encoding β -catenin, leads to ARP2/3 complex overactivity and disordered cortical neuronal migration. <i>Nature Genetics</i> , 2018 , 50, 1093-1101	36.3	48
6	Utility of whole exome sequencing for the early diagnosis of pediatric-onset cerebellar atrophy associated with developmental delay in an inbred population. <i>Orphanet Journal of Rare Diseases</i> , 2016 , 11, 57	4.2	20
5	Inherited glycosylphosphatidylinositol defects cause the rare Emm-negative blood phenotype and developmental disorders. <i>Blood</i> , 2021 , 137, 3660-3669	2.2	7
4	Association of GSTP1 p.Ile105Val (rs1695, c.313A > G) Variant with the Risk of Breast Carcinoma among Egyptian Women. <i>Biochemical Genetics</i> , 2021 , 59, 1487-1505	2.4	1
3	Clinical and genetic characterization of ten Egyptian patients with Wolf-Hirschhorn syndrome and review of literature. <i>Molecular Genetics & Genomic Medicine</i> , 2021 , 9, e1546	2.3	1
2	Serum homocysteine, lipid profile and BMI as atherosclerotic risk factors in children with numerical chromosomal aberrations.. <i>World Journal of Pediatrics</i> , 2022 , 1	4.6	1
1	Clinical Implications of S100A12 and Resolvin D1 Serum Levels, and Related Genes in Children with Familial Mediterranean Fever. <i>Journal of Child Science</i> , 2021 , 11, e163-e169	0.2	