Engy Asem Ashaat

List of Publications by Citations

12 papers	62 citations	4 h-index	7 g-index
14 ext. papers	103 ext. citations	2.2 avg, IF	0.92 L-index

#	Paper	IF	Citations
12	Williams-Beuren syndrome in diverse populations. *American Journal of Medical Genetics, Part A*, **2018**, 176, 1128-1136	2.5	31
11	Turner syndrome in diverse populations. *American Journal of Medical Genetics, Part A*, **2020**, 182, 303-313	2.5	8
10	The potential impact of COMT gene variants on dopamine regulation and phenotypic traits of ASD patients. *Behavioural Brain Research*, **2020**, 378, 112272	3.4	7
9	Study of C677T variant of methylene tetrahydrofolate reductase gene in autistic spectrum disorder Egyptian children. *American Journal of Medical Genetics Part B: Neuropsychiatric Genetics*, **2019**, 180, 305-309	3.5	5
8	Rubinstein-Taybi syndrome in diverse populations. *American Journal of Medical Genetics, Part A*, **2020**, 182, 2939-2950	2.5	4
7	Chromosome 9p terminal deletion in nine Egyptian patients and narrowing of the critical region for trigonocephaly. *Molecular Genetics & Genomic Medicine*, **2021**, 9, e1829	2.3	2
6	A novel homozygous variant in the TRAPPC9 gene causing intellectual disability and autism Spectrum disorder. *Meta Gene*, **2020**, 26, 100783	0.7	1
5	Blepharophimosis-ptosis-intellectual disability syndrome: A report of nine Egyptian patients with further expansion of phenotypic and mutational spectrum. *American Journal of Medical Genetics, Part A*, **2020**, 182, 2857-2866	2.5	1
4	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
3	First Report of Two Egyptian Patients with Desbuquois Dysplasia due to Homozygous Mutations. *Molecular Syndromology*, **2021**, 12, 279-288	1.5	0
2	Assessment of Multiplex Ligation-Dependent Probe Amplification (MLPA) as a diagnostic test for Egyptian patients with Williams-Beuren syndrome. *Gene Reports*, **2020**, 20, 100767	1.4	
1	Phenotypic and Molecular Cytogenetic Analysis of a Case of Monosomy 1p36 Syndrome due to Unbalanced Translocation. *Molecular Syndromology*, **2020**, 11, 284-295	1.5	