Engy Asem Ashaat

List of Publications by Year
in descending order

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14 papers	136 citations	1683354 5 h-index	1281420 11 g-index
14 all docs	14 docs citations	14 times ranked	287 citing authors

#	Article	IF	Citations
1	Williamsâ€“Beuren syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2018, 176, 1128-1136.	0.7	55
2	Rubinsteinâ€“Taybi syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2020, 182, 2939-2950.	0.7	16
3	The potential impact of COMT gene variants on dopamine regulation and phenotypic traits of ASD patients. Behavioural Brain Research, 2020, 378, 112272.	1.2	15
4	Turner syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2020, 182, 303-313.	0.7	15
5	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.	1.1	12
6	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
7	Blepharophimosisâ€ptosisâ€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.	0.7	4
8	Chromosome 9p terminal deletion in nine Egyptian patients and narrowing of the critical region for trigonocephaly. Molecular Genetics & Genomic Medicine, 2021, 9, e1829.	0.6	4
9	First Report of Two Egyptian Patients with Desbuquois Dysplasia due to Homozygous <i>CANT1</i> Mutations. Molecular Syndromology, 2021, 12, 279-288.	0.3	3
10	Altered Adaptive Cellular Immune Function in a Group of Egyptian Children with Autism. Journal of Clinical and Diagnostic Research JCDR, 0, , .	0.8	3
11	A novel homozygous variant in the TRAPPC9 gene causing intellectual disability and autism Spectrum disorder. Meta Gene, 2020, 26, 100783.	0.3	2
12	Phenotypic and Molecular Cytogenetic Analysis of a Case of Monosomy 1p36 Syndrome due to Unbalanced Translocation. Molecular Syndromology, 2020, 11, 284-295.	0.3	1
13	Cover Image, Volume 176A, Number 5, May 2018. American Journal of Medical Genetics, Part A, 2018, 176, .	0.7	0
14	Assessment of Multiplex Ligation-Dependent Probe Amplification (MLPA) as a diagnostic test for Egyptian patients with Williams-Beuren syndrome. Gene Reports, 2020, 20, 100767.	0.4	0