Laila Mahmoud

List of Publications by Year
in descending order

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8 papers	312 citations	1307594 7 h-index	1588992 8 g-index
8 all docs	8 docs citations	8 times ranked	913 citing authors

#	Article	IF	Citations
1	Expanding on the phenotypic spectrum of <scp>Woodhouseâ€Sakati</scp> syndrome due to founder pathogenic variant in <scp>DCAF17</scp>: Report of 58 additional patients from Qatar and literature review. American Journal of Medical Genetics, Part A, 2022, 188, 116-129.	1.2	6
2	Clinical exome sequencing in 509 Middle Eastern families with suspected Mendelian diseases: The Qatari experience. American Journal of Medical Genetics, Part A, 2019, 179, 927-935.	1.2	32
3	Natural history, with clinical, biochemical, and molecular characterization of classical homocystinuria in the Qatari population. Journal of Inherited Metabolic Disease, 2019, 42, 818-830.	3.6	12
4	Bi-allelic TMEM94 Truncating Variants Are Associated with Neurodevelopmental Delay, Congenital Heart Defects, and Distinct Facial Dysmorphism. American Journal of Human Genetics, 2018, 103, 948-967.	6.2	18
5	Clinical genetics and genomic medicine in Qatar. Molecular Genetics & Genomic Medicine, 2018, 6, 702-712.	1.2	12
6	Newborn screening for remethylation disorders and vitamin B12 deficiency-evaluation of new strategies in cohorts from Qatar and Germany. World Journal of Pediatrics, 2017, 13, 136-143.	1.8	24
7	High diagnostic yield of clinical exome sequencing in Middle Eastern patients with Mendelian disorders. Human Genetics, 2015, 134, 967-980.	3.8	168
8	Loss-of-Function Mutation in APC2 Causes Sotos Syndrome Features. Cell Reports, 2015, 10, 1585-1598.	6.4	40