

# Laila Mahmoud

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/11206701/publications.pdf>

Version: 2024-02-01

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