Amal Souissi

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.

13 papers	68 citations	4 h-index	8 g-index
16 ext. papers	158 ext. citations	5.8 avg, IF	1.7 L-index

#	Paper	IF	Citations
13	A year of genomic surveillance reveals how the SARS-CoV-2 pandemic unfolded in Africa. *Science*, **2021**, 374, 423-431	33.3	35
12	Novel pathogenic mutations and further evidence for clinical relevance of genes and variants causing hearing impairment in Tunisian population. *Journal of Advanced Research*, **2021**, 31, 13-24	13	8
11	SRD5A3-CDG: 3D structure modeling, clinical spectrum, and computer-based dysmorphic facial recognition. *American Journal of Medical Genetics, Part A*, **2021**, 185, 1081-1090	2.5	8
10	Evidence of SARS-CoV-2 Symptomatic Reinfection in Four Health Care Professionals from the Same Hospital Despite the Presence of Antibodies.. *International Journal of Infectious Diseases*, **2022**,	10.5	4
9	Osteoprotegerin gene polymorphisms and otosclerosis: an additional genetic association study, multilocus interaction and meta-analysis. *BMC Medical Genetics*, **2020**, 21, 122	2.1	3
8	A year of genomic surveillance reveals how the SARS-CoV-2 pandemic unfolded in Africa		3
7	8q21.11 microdeletion syndrome: Delineation of HEY1 as a candidate gene in neurodevelopmental and cardiac defects. *Molecular Genetics & Genomic Medicine*, **2021**, 9, e1811	2.3	2
6	Custom Next-Generation Sequencing Identifies Novel Mutations Expanding the Molecular and clinical spectrum of isolated Hearing Impairment or along with defects of the retina, the thyroid, and the kidneys.. *Molecular Genetics & Genomic Medicine*, **2022**, e1868	2.3	1
5	Genetics and meta-analysis of recessive non-syndromic hearing impairment and Usher syndrome in Maghreb population: lessons from the past, contemporary actualities and future challenges. *Human Genetics*, **2021**, 1	6.3	1
4	Molecular insights into kinase domain variants explain variability in both severity and progression of DFNB30 hearing impairment. *Journal of Biomolecular Structure and Dynamics*, **2021**, 1-12	3.6	1
3	Expanding the Clinical and Molecular Spectrum of HARS2-Perrault Syndrome: Identification of a Novel Homozygous Missense Variant in the gene. *Genetic Testing and Molecular Biomarkers*, **2021**, 25, 528-539	1.6	1
2	Further insights into the spectrum phenotype of TRAPPC9 and CDK5RAP2 genes, segregating independently in a large Tunisian family with intellectual disability and microcephaly. *European Journal of Medical Genetics*, **2021**, 64, 104373	2.6	
1	Gene duplication and functional divergence of the zebrafish otospiralin genes. *Development Genes and Evolution*, **2020**, 230, 27-36	1.8	