S Zeinali

List of Publications by Year
in descending order

12 papers	261 citations	1477746 6 h-index	1199166 12 g-index
12 all docs	12 docs citations	12 times ranked	762 citing authors

#	Article	IF	Citations
1	Mapping autosomal recessive intellectual disability: combined microarray and exome sequencing identifies 26 novel candidate genes in 192 consanguineous families. Molecular Psychiatry, 2018, 23, 973-984.	4.1	147
2	Autosomal recessive congenital ichthyosis: Genomic landscape and phenotypic spectrum in a cohort of 125 consanguineous families. Human Mutation, 2019, 40, 288-298.	1.1	43
3	Lessons for preparedness and reasons for concern from the early COVID-19 epidemic in Iran. Epidemics, 2021, 36, 100472.	1.5	17
4	Phenotypic heterogeneity inPIK3CA-related overgrowth spectrum. British Journal of Dermatology, 2016, 175, 810-814.	1.4	10
5	Molecular genetic diagnosis of Glanzmann syndrome in Iranian population; reporting novel and recurrent mutations. Orphanet Journal of Rare Diseases, 2019, 14, 87.	1.2	10
6	Erythrokeratoderma: a manifestation associated with multiple types of ichthyoses with different gene defects. British Journal of Dermatology, 2018, 178, e219-e221.	1.4	7
7	Association of xenobiotic-metabolizing enzymes (GSTM1 and GSTT 1), and pro-inflammatory cytokines (TNF-Î± and IL-6) genetic polymorphisms with non-alcoholic fatty liver disease. Molecular Biology Reports, 2021, 48, 1225-1231.	1.0	7
8	Targeted integration into pseudo attP sites of CHO cells using CRISPR/Cas9. Journal of Biotechnology, 2021, 337, 1-7.	1.9	6
9	Frequencies of intron 1 and 22 inversions of factor <scp>VIII</scp> gene: A first report in Afghan patients with severe haemophilia A. Haemophilia, 2018, 24, e157-e160.	1.0	5
10	Prevalence of factor VIII inhibitors among Afghan patients with hemophilia A. Blood Coagulation and Fibrinolysis, 2018, 29, 697-700.	0.5	5
11	Ichthyosis, psoriasiform dermatitis, and recurrent fungal infections in patients with biallelic mutations in PERP. Journal of the European Academy of Dermatology and Venereology, 2022, 36, 472-479.	1.3	3
12	Development and implementation of a novel panel consisting 20 markers for the detection of genetic causes of male infertility. Andrologia, 2018, 50, e12946.	1.0	1