

S Zeinali

List of Publications by Year in descending order

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Version: 2024-02-01

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	Ichthyosis, psoriasiform dermatitis, and recurrent fungal infections in patients with biallelic mutations in <i>PERP</i> . <i>Journal of the European Academy of Dermatology and Venereology</i> , 2022, 36, 472-479.	1.3	3
2	Targeted integration into pseudo attP sites of CHO cells using CRISPR/Cas9. <i>Journal of Biotechnology</i> , 2021, 337, 1-7.	1.9	6
3	Lessons for preparedness and reasons for concern from the early COVID-19 epidemic in Iran. <i>Epidemics</i> , 2021, 36, 100472.	1.5	17
4	Association of xenobiotic-metabolizing enzymes (GSTM1 and GSTT 1), and pro-inflammatory cytokines (TNF- α and IL-6) genetic polymorphisms with non-alcoholic fatty liver disease. <i>Molecular Biology Reports</i> , 2021, 48, 1225-1231.	1.0	7
5	Molecular genetic diagnosis of Glanzmann syndrome in Iranian population; reporting novel and recurrent mutations. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 87.	1.2	10
6	Autosomal recessive congenital ichthyosis: Genomic landscape and phenotypic spectrum in a cohort of 125 consanguineous families. <i>Human Mutation</i> , 2019, 40, 288-298.	1.1	43
7	Frequencies of intron 1 and 22 inversions of factor <i>VIII</i> gene: A first report in Afghan patients with severe haemophilia A. <i>Haemophilia</i> , 2018, 24, e157-e160.	1.0	5
8	Development and implementation of a novel panel consisting 20 markers for the detection of genetic causes of male infertility. <i>Andrologia</i> , 2018, 50, e12946.	1.0	1
9	Mapping autosomal recessive intellectual disability: combined microarray and exome sequencing identifies 26 novel candidate genes in 192 consanguineous families. <i>Molecular Psychiatry</i> , 2018, 23, 973-984.	4.1	147
10	Erythrokeratoderma: a manifestation associated with multiple types of ichthyoses with different gene defects. <i>British Journal of Dermatology</i> , 2018, 178, e219-e221.	1.4	7
11	Prevalence of factor VIII inhibitors among Afghan patients with hemophilia A. <i>Blood Coagulation and Fibrinolysis</i> , 2018, 29, 697-700.	0.5	5
12	Phenotypic heterogeneity in PIK3CA-related overgrowth spectrum. <i>British Journal of Dermatology</i> , 2016, 175, 810-814.	1.4	10