Marzieh Mojbafan

List of Publications by Year
in descending order

15	121	1477746 6	1281420 11
papers	citations	h-index	g-index

16	16	16	205
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The expression of peroxisomal protein transcripts increased by retinoic acid during neural differentiationâ˜†. Differentiation, 2011, 81, 127-132.	1.0	40
2	In silico analysis of novel mutations in maple syrup urine disease patients from Iran. Metabolic Brain Disease, 2017, 32, 105-113.	1.4	16
3	Mutational spectrum of autosomal recessive limb-girdle muscular dystrophies in a cohort of 112 Iranian patients and reporting of a possible founder effect. Orphanet Journal of Rare Diseases, 2020, 15, 14.	1.2	13
4	Reporting a rare form of myopathy, myopathy with extrapyramidal signs, in an Iranian family using next generation sequencing: a case report. BMC Medical Genetics, 2020, 21, 77.	2.1	11
5	Hints From the Cellular Functions to the Practical Outlook of Circular RNAs. Frontiers in Genetics, 2021, 12, 679446.	1.1	9
6	Sex-specific association of RANTES gene â˚'403 variant in Menièreâ€™s disease. European Archives of Oto-Rhino-Laryngology, 2015, 272, 2221-2225.	0.8	6
7	Liver alpha-amylase gene expression as an early obesity biomarker. Pharmacological Reports, 2017, 69, 229-234.	1.5	6
8	Linkage Study Revealed Complex Haplotypes in a Multifamily due to Different Mutations in CAPN3 Gene in an Iranian Ethnic Group. Journal of Molecular Neuroscience, 2016, 59, 392-396.	1.1	5
9	A rare form of limb girdle muscular dystrophy (type 2E) seen in an Iranian family detected by autozygosity mapping. Journal of Neurogenetics, 2016, 30, 1-4.	0.6	4
10	Molecular genetic study of Calpainopathy in Iran. Gene, 2018, 677, 259-265.	1.0	4
11	A novel mutation in alpha sarcoglycan gene in an Iranian family with limb girdle muscular dystrophy 2D. Neurological Research, 2016, 38, 220-223.	0.6	2
12	Genetic variability in Iranian limbâ€girdle muscular dystrophy type 2B patients: An evidence of a founder effect. Molecular Genetics & Genomic Medicine, 2019, 7, e1029.	0.6	2
13	Identification of a novel mutation in congenital afibrinogenemia in Iranian patients. Blood Coagulation and Fibrinolysis, 2021, 32, 323-327.	0.5	1
14	Genetic study of a patient with congenital central hypoventilation syndrome in Iran: a case report. Molecular Biology Reports, 2021, 48, 8239-8243.	1.0	1
15	A decade of molecular preimplantation genetic diagnosis of 350 blastomeres for beta-thalassemia combined with HLA typing, aneuploidy screening and sex selection in Iran. BMC Pregnancy and Childbirth, 2022, 22, 330.	0.9	1