

Marzieh Mojbafan

List of Publications by Year in descending order

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15
papers

121
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1477746

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1281420

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205
citing authors

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