Kolsoum Inanloorahatloo

List of Publications by Year
in descending order

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16 papers	324 citations	1307594 7 h-index	1199594 12 g-index
16 all docs	16 docs citations	16 times ranked	844 citing authors

#	Article	IF	Citations
1	Intellectual disability associated with craniofacial dysmorphism due to POLR3B mutation and defect in spliceosomal machinery. BMC Medical Genomics, 2022, 15, 89.	1.5	1
2	Intermittent white noise exposure is associated with rat cochleae damage and changes in the gene expression. Egyptian Journal of Medical Human Genetics, 2022, 23, .	1.0	0
3	Evolution of SARSâ€CoVâ€2 genome from December 2019 to late March 2020: EmergedÂhaplotypes and informative TagÂnucleotide variations. Journal of Medical Virology, 2021, 93, 2010-2020.	5.0	10
4	Deep geno- and phenotyping in two consanguineous families with CMT2 reveals HADHA as an unusual disease-causing gene and an intronic variant in GDAP1 as an unusual mutation. Journal of Neurology, 2021, 268, 640-650.	3.6	2
5	Worldâ€wide tracking of major SARSâ€CoVâ€2 genome haplotypes in sequences of June 1 to November 15, 2020 and discovery of rapid expansion of a new haplotype. Journal of Medical Virology, 2021, 93, 3251-3256.	5.0	4
6	Study on SARS-CoV-2 strains in Iran reveals potential contribution of co-infection with and recombination between different strains to the emergence of new strains. Virology, 2021, 562, 63-73.	2.4	26
7	Whole-Transcriptome Analysis Reveals Dysregulation of Actin-Cytoskeleton Pathway in Intellectual Disability Patients. Neuroscience, 2019, 404, 423-444.	2.3	4
8	Sex-based differences in myocardial gene expression in recently deceased organ donors with no prior cardiovascular disease. PLoS ONE, 2017, 12, e0183874.	2.5	43
9	iPSC-derived cardiomyocytes reveal abnormal TGF-Î² signalling in left ventricular non-compaction cardiomyopathy. Nature Cell Biology, 2016, 18, 1031-1042.	10.3	148
10	Polymorphisms of cystathionine beta-synthase gene are associated with susceptibility to sepsis. European Journal of Human Genetics, 2016, 24, 1041-1048.	2.8	8
11	A Rapid, High-Quality, Cost-Effective, Comprehensive and Expandable Targeted Next-Generation Sequencing Assay for Inherited Heart Diseases. Circulation Research, 2015, 117, 603-611.	4.5	34
12	Abstract 248: Aberrant TGFÎ² Signaling as an Etiology of Left Ventricular Non-compaction Cardiomyopathy. Circulation Research, 2015, 117, .	4.5	0
13	Mutation in ST6GALNAC5 identified in family with coronary artery disease. Scientific Reports, 2014, 4, 3595.	3.3	23
14	Mutation in CYP27A1 identified in family with coronary artery disease. European Journal of Medical Genetics, 2013, 56, 655-660.	1.3	15
15	Expression of Dmrt Family Genes During Gonadal Differentiation in Two Species of Artemia (Branchiopoda, Anostraca) from Urmia Lake (Iran). Crustaceana, 2010, 83, 1153-1165.	0.3	6
16	Comparison of the protein profile of the reproductive system between bisexual and parthenogenetic species of Artemia (Branchiopoda, Anostraca). Crustaceana, 2009, 82, 1237-1248.	0.3	0