Morteza oladnabi

List of Publications by Year
in descending order

29 papers	468 citations	933447 10 h-index	713466 21 g-index
29 all docs	29 docs citations	29 times ranked	1174 citing authors

#	Article	IF	Citations
1	Molecular insight of dyskeratosis congenita: Defects in telomere length homeostasis.. Journal of Clinical and Translational Research, 2022, 8, 20-30.	0.3	0
2	Correlation between ELFâ€“PEMF exposure and Human RPE Cell Proliferation, Apoptosis and Gene Expression. Journal of Ophthalmic and Vision Research, 2021, 16, 202-211.	1.0	0
3	WDR81 Gene Silencing Can Reduce Exosome Levels in Human U87-MG GlioblastomaÂCells. Journal of Molecular Neuroscience, 2021, 71, 1696-1702.	2.3	3
4	In silico drug repurposing for the treatment of heart diseases using gene expression data and molecular docking techniques. Biochemical and Biophysical Research Communications, 2021, 572, 138-144.	2.1	1
5	Cannabinoid CB2 Receptor Functional Variation (Q63R) Is Associated with Multiple Sclerosis in Iranian Subjects. Journal of Molecular Neuroscience, 2020, 70, 26-31.	2.3	11
6	Molecular and biochemical mechanisms of human iris color: A comprehensive review. Journal of Cellular Physiology, 2020, 235, 8972-8982.	4.1	13
7	Variants in Intron 4 of PD-1 Gene are Associated with the Susceptibility to SLE in an Iranian Population. Iranian Journal of Immunology, 2020, 17, 204-214.	0.6	0
8	Toll-Like Receptor (TLR)-9 rs352140 Polymorphism is an Immunopathology Protective Factor in Parkinson's Disease in the Northern Iranian Population. Iranian Journal of Immunology, 2020, 17, 313-323.	0.6	1
9	Iranome: A catalog of genomic variations in the Iranian population. Human Mutation, 2019, 40, 1968-1984.	2.5	116
10	Distinct genetic variation and heterogeneity of the Iranian population. PLoS Genetics, 2019, 15, e1008385.	3.5	34
11	Evaluation of cytokeratin 19 as a prognostic tumoral and metastatic marker with focus on improved detection methods. Journal of Cellular Physiology, 2019, 234, 21425-21435.	4.1	21
12	Genetics of intellectual disability in consanguineous families. Molecular Psychiatry, 2019, 24, 1027-1039.	7.9	131
13	Extremely low frequency-pulsed electromagnetic fields affect proangiogenic-related gene expression in retinal pigment epithelial cells. Iranian Journal of Basic Medical Sciences, 2019, 22, 128-133.	1.0	5
14	Myeloid Cell Leukemia-1 (MCL-1) siRNA Therapy Showed Cytotoxic Effect on T Cells Acute Lymphoblastic Leukemia. International Journal of Cancer Management, 2019, In Press, .	0.4	0
15	Enzymatic characterization of a NADH-dependent diaphorase from Lysinibacillus sp. strain PAD-91. Protein Expression and Purification, 2018, 146, 1-7.	1.3	2
16	CNKSR1 gene defect can cause syndromic autosomal recessive intellectual disability. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2018, 177, 691-699.	1.7	4
17	Gene Silencing of TGFÎ²RII Can Inhibit Glioblastoma Cell Growth. Asian Pacific Journal of Cancer Prevention, 2018, 19, 2681-2686.	1.2	3
18	Recombinant expression, characterization and application of a dihydrolipoamide dehydrogenase with diaphorase activity from Bacillus sphaericus. 3 Biotech, 2017, 7, 153.	2.2	3

#	Article	IF	Citations
19	Purification and Characterization of Recombinant Darbepoetin Alfa from Leishmania tarentolae. Molecular Biotechnology, 2016, 58, 566-572.	2.4	1
20	Cloning and expression of codon-optimized recombinant darbepoetin alfa in Leishmania tarentolae T7-TR. Protein Expression and Purification, 2016, 118, 120-125.	1.3	7
21	The Rapid and Sensitive Quantitative Determination of Galactose by Combined Enzymatic and Colorimetric Method: Application in Neonatal Screening. Applied Biochemistry and Biotechnology, 2016, 179, 283-293.	2.9	4
22	New evidence for the role of calpain 10 in autosomal recessive intellectual disability: identification of two novel nonsense variants by exome sequencing in Iranian families. Archives of Iranian Medicine, 2015, 18, 179-84.	0.6	6
23	Aberrant expression of Activating Transcription Factor 6 (ATF6) in major psychiatric disorders. Psychiatry Research, 2012, 200, 1086-1087.	3.3	3
24	Novel evidence of the involvement of calreticulin in major psychiatric disorders. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2012, 37, 276-281.	4.8	16
25	Core promoter STRs: Novel mechanism for inter-individual variation in gene expression in humans. Gene, 2012, 492, 195-198.	2.2	25
26	Exceptional human core promoter nucleotide compositions. Gene, 2011, 475, 79-86.	2.2	19
27	Reversion of the human calreticulin gene promoter to the ancestral type as a result of a novel psychosis-associated mutation. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2011, 35, 541-544.	4.8	18
28	Novel extreme homozygote haplotypes at the human caveolin 1 gene upstream purine complex in sporadic Alzheimer's disease. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2010, 153B, 347-349.	1.7	11
29	Novel mutations in the calreticulin gene core promoter and coding sequence in schizoaffective disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2010, 153B, 706-709.	1.7	10