Reza Alibakhshi

List of Publications by Year
in descending order

35	319	932766 10	887659 17
papers	citations	h-index	g-index

35	35	35	430
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Î±-Thalassemia Mutations in Ilam Province, West Iran. Hemoglobin, 2022, 46, 147-152.	0.4	6
2	Spectrum of MEFV gene mutations in 4,256 familial Mediterranean fever patients from Iran: a comprehensive systematic review. Egyptian Journal of Medical Human Genetics, 2022, 23, .	0.5	2
3	The pathogenicity classification of PAH gene variants in the Iranian population. Computational Biology and Chemistry, 2022, 98, 107665.	1.1	1
4	Problem of borderline hemoglobin A2 levels in an Iranian population with a high prevalence of Î±- and Î²-thalassemia carriers. Egyptian Journal of Medical Human Genetics, 2022, 23, .	0.5	1
5	CFTR gene mutation spectrum among 735 Iranian patients with cystic fibrosis: A comprehensive systematic review. Pediatric Pulmonology, 2021, 56, 3644-3656.	1.0	5
6	Spectrum of PAH gene mutations in 1547 phenylketonuria patients from Iran: a comprehensive systematic review. Metabolic Brain Disease, 2021, 36, 767-780.	1.4	7
7	Distribution of HBB Gene Mutations in the Kurdish Population of Ilam Province, West Iran. Hemoglobin, 2020, 44, 244-248.	0.4	3
8	Molecular Genetic Analysis of Î±-Thalassemia in Hamadan Province, West Iran. Hemoglobin, 2020, 44, 319-324.	0.4	4
9	Severe Î±-Thalassemia Due to Compound Heterozygosity for Hb Adana (Î±59 Gly>Asp) (HBA1: c.179Gâ€‰>â€‰A) and Codon 127 (Aâ€‰>â€‰T) (HBA2: c.382Aâ€‰>â€‰T) in an Iranian Family. Hemoglobin, 2020, 44, 139-142.	0.4	0
10	Haplotype analysis of the CFTR gene on normal and mutant CFTR genes. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2020, 821, 111708.	0.4	2
11	The Spectrum of Î±-Thalassemia Mutations in Kurdistan Province, West Iran. Hemoglobin, 2020, 44, 156-161.	0.4	6
12	Cytogenetic analysis of 570 couples with recurrent pregnancy loss: Reporting 11 years of experience. Journal of Human Reproductive Sciences, 2020, 13, 216.	0.4	4
13	The Spectrum of Î±-Thalassemia Mutations in the Lak Population of Iran. Hemoglobin, 2019, 43, 107-111.	0.4	11
14	The Spectrum of Î²-Thalassemia Mutations in Hamadan Province, West Iran. Hemoglobin, 2019, 43, 18-22.	0.4	11
15	Characterization of the IVS-II-821 (A>C) (HBB: c.316-30A>C) Mutation in a Î²-Thalassemia Phenotype in Iran. Hemoglobin, 2019, 43, 23-26.	0.4	0
16	Association between activity and genotypes of paraoxonase1 L55M (rs854560) increases the disease activity of rheumatoid arthritis through oxidative stress. Molecular Biology Reports, 2019, 46, 741-749.	1.0	8
17	The status of gene-VNTR alleles and mini-haplotypes associations with gene mutations in Iranian Kurdish PKU patients. Medical Journal of the Islamic Republic of Iran, 2019, 33, 88.	0.9	0
18	Mutation Analysis in Western Iran: Identification of Two Novel Mutations. Journal of Reproduction and Infertility, 2018, 19, 3-9.	1.0	7

#	Article	IF	Citations
19	Spectrum of Phenylalanine Hydroxylase Gene Mutations in Hamadan and Lorestan Provinces of Iran and Their Associations with Variable Number of Tandem Repeat Alleles. Iranian Journal of Medical Sciences, 2018, 43, 318-323.	0.3	9
20	Detection of SPG20 gene promoter-methylated DNA, as a novel epigenetic biomarker, in plasma for colorectal cancer diagnosis using the MethyLight method. Oncology Letters, 2017, 13, 3277-3284.	0.8	33
21	Molecular analysis of exon 13 of cystic fibrosis patients in Middle East: High frequency of K710X mutation. Meta Gene, 2017, 11, 123-126.	0.3	0
22	IVS-II-648/649 (â€“T) (HBB: c.316â^'202del) Triggers a Novel Î²-Thalassemia Phenotype. Hemoglobin, 2017, 41, 44-46.	0.4	3
23	TTY2 genes deletions as genetic risk factor of male infertility. Cellular and Molecular Biology, 2017, 63, 57.	0.3	3
24	Aberrant Methylation of APAF-1 Gene in Acute Myeloid Leukemia Patients. International Journal of Hematology-Oncology and Stem Cell Research, 2017, 11, 225-230.	0.3	2
25	Hepatitis C Virus Genotype Distribution in Kermanshah Province, Western Iran. Modern Applied Science, 2015, 10, 138.	0.4	0
26	The Spectrum ofÎ±-Thalassemia Mutations in Kermanshah Province, West Iran. Hemoglobin, 2015, 39, 403-406.	0.4	16
27	Mutation analysis of PAH gene in patients with PKU in western Iran and its association with polymorphisms: identification of four novel mutations. Metabolic Brain Disease, 2014, 29, 131-138.	1.4	31
28	Mutation Analysis of Exons 10 and 17a of CFTR Gene in Patients with Cystic Fibrosis in Kermanshah Province, Western Iran. Journal of Reproduction and Infertility, 2014, 15, 49-56.	1.0	5
29	The Spectrum of Î²-Thalassemia Mutations in Kermanshah Province in West Iran and its Association with Hematological Parameters. Hemoglobin, 2013, 37, 544-552.	0.4	12
30	The proportion of tetrahydrobiopterin deficiency and PAH gene deficiency variants among cases with hyperphenyalaninemia in Western Iran. Indian Journal of Human Genetics, 2013, 19, 454.	0.7	4
31	Molecular analysis of exons 6 and 7 of phenylalanine hydroxylase gene mutations in Phenylketonuria patients in Western Iran. Indian Journal of Human Genetics, 2012, 18, 290.	0.7	16
32	Paraoxonase Arg 192 allele is an independent risk factor for three-vessel stenosis of coronary artery disease. Molecular Biology Reports, 2011, 38, 5421-5428.	1.0	24
33	Analysis of the CFTR gene in Iranian cystic fibrosis patients: Identification of eight novel mutations. Journal of Cystic Fibrosis, 2008, 7, 102-109.	0.3	63
34	Mutation analysis of CFTR gene in 70 Iranian cystic fibrosis patients. Iranian Journal of Allergy, Asthma and Immunology, 2006, 5, 3-8.	0.3	20
35	The status of PAH gene-VNTR alleles and mini-haplotypes associations with PAH gene mutations in Iranian Kurdish PKU patients. Medical Journal of the Islamic Republic of Iran, 0, , .	0.9	0