

Maryam Neishabury

List of Publications by Year in descending order

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18
papers

270
citations

933447

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996975

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docs citations

18
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377
citing authors

#	ARTICLE	IF	CITATIONS
1	Spontaneous mutation, oxidative DNA damage, and the roles of base and nucleotide excision repair in the yeast <i>Saccharomyces cerevisiae</i> . <i>Yeast</i> , 1999, 15, 205-218.	1.7	57
2	The XmnI and BCL11A Single Nucleotide Polymorphisms May Help Predict Hydroxyurea Response in Iranian β^2 -Thalassemia Patients. <i>Hemoglobin</i> , 2012, 36, 371-380.	0.8	35
3	Rare and unexpected mutations among Iranian beta-thalassemia patients and prenatal samples discovered by reverse-hybridization and DNA sequencing. <i>Haematologica</i> , 2002, 87, 1113-4.	3.5	26
4	The Iranian Human Mutation Gene Bank: A data and sample resource for worldwide collaborative genetics research. <i>Human Mutation</i> , 2003, 21, 146-150.	2.5	25
5	High Prevalence of the $\Delta 3.7$ Deletion Among Thalassemia Patients in Iran. <i>Hemoglobin</i> , 2003, 27, 53-55.	0.8	21
6	Molecular Mechanisms Underlying Thalassemia Intermedia in Iran. <i>Genetic Testing and Molecular Biomarkers</i> , 2008, 12, 549-556.	1.7	21
7	Frequency of Positive XmnI G ³ polymorphism and coinheritance of common alpha thalassemia mutations do not show statistically significant difference between thalassemia major and intermedia cases with homozygous IVSII-1 mutation. <i>Blood Cells, Molecules, and Diseases</i> , 2010, 44, 95-99.	1.4	20
8	alpha-globin gene deletion and point mutation analysis among in Iranian patients with microcytic hypochromic anemia. <i>Haematologica</i> , 2003, 88, 1196-7.	3.5	19
9	The influence of the BCL11A polymorphism on the phenotype of patients with beta thalassemia could be affected by the beta globin locus control region and/or the XmnI-HBG2 genotypic background. <i>Blood Cells, Molecules, and Diseases</i> , 2013, 51, 80-84.	1.4	11
10	The modifying effect of XmnI-HBG2 on thalassemic phenotype is associated with its linked elements in the beta globin locus control region, including the palindromic site at 5 ² HS4. <i>Blood Cells, Molecules, and Diseases</i> , 2012, 48, 1-5.	1.4	10
11	Novel mutations in mitochondrial carrier family gene SLC25A38, causing congenital sideroblastic anemia in Iranian families, identified by whole exome sequencing. <i>Blood Cells, Molecules, and Diseases</i> , 2018, 71, 39-44.	1.4	7
12	Commentary on "The modifying effect of XmnI-HBG2 on thalassemic phenotype is associated with its linked elements in the beta globin locus control region, including the palindromic site at 5 ² HS4" by M. Neishabury et al.. <i>Blood Cells, Molecules, and Diseases</i> , 2012, 48, 6.	1.4	5
13	A REPORT OF 8 CASES WITH HEMOGLOBIN H DISEASE IN AN IRANIAN FAMILY. <i>Pediatric Hematology and Oncology</i> , 2010, 27, 405-412.	0.8	4
14	Analyzing 5 ² HS3 and 5 ² HS4 LCR core regions and NF-E2 in Iranian thalassemia intermedia patients with normal or carrier status for beta-globin mutations. <i>Blood Cells, Molecules, and Diseases</i> , 2011, 46, 201-205.	1.4	3
15	Novel variants in Iranian individuals suspected to have inherited red blood cell disorders, including bone marrow failure syndromes. <i>Haematologica</i> , 2020, 105, e1-e4.	3.5	3
16	The First Case of BENTA Disease (B Cell Expansion with NF- β B and T Cell Anergy) from Iran. <i>Journal of Clinical Immunology</i> , 2021, 41, 811-813.	3.8	2
17	Electromobility Shift Assay Reveals Evidence in Favor of Allele-Specific Binding of RUNX1 to the 5 ² Hypersensitive Site 4-Locus Control Region. <i>Hemoglobin</i> , 2016, 40, 236-239.	0.8	1
18	The Impact of XmnI-HBG2, BCL11A and HBS1L-MYB Single Nucleotide Polymorphisms on Hb F Variation of Hematologically Normal Iranian Individuals. <i>Hemoglobin</i> , 2016, 40, 198-201.	0.8	0