

Mohammad Keramatipour

List of Publications by Year in descending order

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Version: 2024-02-01

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papers

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citations

1040056

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996975

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docs citations

41
times ranked

737
citing authors

#	ARTICLE	IF	CITATIONS
1	Effect of DHA-rich fish oil on PPAR α target genes related to lipid metabolism in type 2 diabetes: A randomized, double-blind, placebo-controlled clinical trial. <i>Journal of Clinical Lipidology</i> , 2015, 9, 770-777.	1.5	43
2	Dysfunction of the ciliary ARMC9/TOGARAM1 protein module causes Joubert syndrome. <i>Journal of Clinical Investigation</i> , 2020, 130, 4423-4439.	8.2	43
3	The importance of CDC27 in cancer: molecular pathology and clinical aspects. <i>Cancer Cell International</i> , 2021, 21, 160.	4.1	24
4	APOA II genotypes frequency and their interaction with saturated fatty acids consumption on lipid profile of patients with type 2 diabetes. <i>Clinical Nutrition</i> , 2016, 35, 907-911.	5.0	15
5	The association between <sc>ST</sc> 18 gene polymorphism and severe pemphigus disease among Iranian population. <i>Experimental Dermatology</i> , 2018, 27, 1395-1398.	2.9	15
6	Genetic mutations and immunological features of severe combined immunodeficiency patients in Iran. <i>Immunology Letters</i> , 2019, 216, 70-78.	2.5	14
7	Genetic study of the PAH locus in the Iranian population: familial gene mutations and minihaplotypes. <i>Metabolic Brain Disease</i> , 2017, 32, 1685-1691.	2.9	13
8	Multi affected pedigree with congenital microcephaly: WES revealed PNKP gene mutation. <i>Brain and Development</i> , 2019, 41, 182-186.	1.1	11
9	Comparison of the effects of nobiletin and letrozole on the activity and expression of aromatase in the MCF-7 breast cancer cell line. <i>Biochemistry and Cell Biology</i> , 2017, 95, 468-473.	2.0	10
10	A Case Series: Congenital Hyperinsulinism. <i>International Journal of Endocrinology and Metabolism</i> , 2016, 14, e37311.	1.0	9
11	Whole Exome Sequencing Reveals a BSCL2 Mutation Causing Progressive Encephalopathy with Lipodystrophy (PELD) in an Iranian Pediatric Patient. <i>Iranian Biomedical Journal</i> , 2016, 20, 295-301.	0.7	8
12	Targeted next generation sequencing identified a novel mutation in MYO7A causing Usher syndrome type 1 in an Iranian consanguineous pedigree. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2018, 104, 10-13.	1.0	7
13	Homozygosity Mapping and Targeted Sanger Sequencing Identifies Three Novel CRB1 (Cumbs) Tj ETQq1 1 0.784314 rgBT /Overlock 294-302.	0.7	7
14	Single Nucleotide Polymorphism rs 2476601 of PTPN22 Gene and Susceptibility to Rheumatoid Arthritis in Iranian Population. <i>Iranian Journal of Allergy, Asthma and Immunology</i> , 2015, 14, 437-42.	0.4	7
15	A Novel Mutation in Gene Causes Malignant Infantile Osteopetrosis. <i>Avicenna Journal of Medical Biotechnology</i> , 2017, 9, 205-208.	0.3	6
16	Effect of Whole Exome Sequencing in Diagnosis of Inborn Errors of Metabolism and Neurogenetic Disorders. <i>Iranian Journal of Child Neurology</i> , 2018, 12, 7-15.	0.3	6
17	First report of Klein-Waardenburg Syndrome in Iran and a novel pathogenic splice site variant in PAX3 gene. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2018, 113, 229-233.	1.0	5
18	Exome sequencing reveals novel rare variants in Iranian familial multiple sclerosis: The importance of POLD2 in the disease pathogenesis. <i>Genomics</i> , 2021, 113, 2645-2655.	2.9	5

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19	Three Novel Variants identified in <i>FBN1</i> and <i>TGFBR2</i> in seven Iranian families with suspected Marfan syndrome. <i>Molecular Genetics & Genomic Medicine</i> , 2020, 8, e1274.	1.2	4
20	Four mutations in <i>MITF</i> , <i>SOX10</i> and <i>PAX3</i> genes were identified as genetic causes of waardenburg syndrome in four unrelated Iranian patients: case report. <i>BMC Pediatrics</i> , 2021, 21, 70.	1.7	4
21	Whole Exome Sequencing Reveals a <i>XPNPEP3</i> Novel Mutation Causing Nephronophthisis in a Pediatric Patient. <i>Iranian Biomedical Journal</i> , 2020, 24, 400-403.	0.7	4
22	Identification of a Novel Mutation in the <i>PAH</i> Gene in an Iranian Phenylketonuria Family: A Case Report. <i>Iranian Journal of Public Health</i> , 2017, 46, 560-564.	0.5	4
23	SPOAN syndrome: a novel mutation and new ocular findings; a case report. <i>BMC Neurology</i> , 2021, 21, 24.	1.8	3
24	Identification of a novel truncating variant in <i>AHL1</i> gene and a brief review on mutations spectrum. <i>Molecular Biology Reports</i> , 2021, 48, 5339-5345.	2.3	3
25	X-linked SCID with a rare mutation. <i>Allergy, Asthma and Clinical Immunology</i> , 2021, 17, 107.	2.0	3
26	Whole-Exome Sequencing Identified a Novel Variant (C.405_422+39del) in <i>DSP</i> Gene in an Iranian Pedigree with Familial Dilated Cardiomyopathy. <i>Reports of Biochemistry and Molecular Biology</i> , 2021, 10, 280-287.	1.4	2
27	Once in a Blue Moon, a Very Rare Coexistence of Glutaric Acidemia Type I and Mucopolysaccharidosis Type IIIB in a Patient. <i>Iranian Biomedical Journal</i> , 2020, 24, 201-205.	0.7	2
28	An Efficient Trio-Based Mini-Haplotyping Method for Genetic Diagnosis of Phenylketonuria. <i>Cell Journal</i> , 2016, 18, 229-36.	0.2	2
29	Comparison of Two Different PCR-based Methods for Detection of GAA Expansions in Frataxin Gene. <i>Iranian Journal of Public Health</i> , 2017, 46, 222-228.	0.5	2
30	A Novel Variant in the Gene Causing Phenylketonuria in an Iranian Pedigree. <i>Avicenna Journal of Medical Biotechnology</i> , 2017, 9, 146-149.	0.3	2
31	Investigating the Effect of rs3783605 Single-nucleotide Polymorphism on the Activity of VCAM-1 Promoter in Human Umbilical Vein Endothelial Cells. <i>Iranian Journal of Allergy, Asthma and Immunology</i> , 2015, 14, 179-87.	0.4	2
32	PKD2 mutation in an Iranian autosomal dominant polycystic kidney disease family with misleading linkage analysis data. <i>Kidney Research and Clinical Practice</i> , 2016, 35, 96-101.	2.2	1
33	Identifying Novel Mutations in Iranian Patients with LPS-responsive Beige-like Anchor Protein (LRBA) Deficiency. <i>Immunological Investigations</i> , 2021, 50, 399-405.	2.0	1
34	The Relationship Between Coronary Artery Disease and Genetic Polymorphisms of Melanoma Inhibitory Activity 3. <i>Iranian Red Crescent Medical Journal</i> , 2016, 18, e31146.	0.5	1
35	Investigation of the association of G-7A and T-138C single nucleotide polymorphisms on the promoter of <i>MGP</i> gene with renal stone. <i>Journal of Nephro pharmacology</i> , 2018, 7, 145-148.	0.4	1
36	Functional Analysis of A Novel Splicing Mutation in The Mutase Gene of Two Unrelated Pedigrees. <i>Cell Journal</i> , 2016, 18, 397-404.	0.2	1

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37	CRB1-Related Leber Congenital Amaurosis: Reporting Novel Pathogenic Variants and a Brief Review on Mutations Spectrum. Iranian Biomedical Journal, 2019, 23, 362-8.	0.7	1
38	Succinate Dehydrogenase Deficiency: A Treatable Neurometabolic Disorder. Iranian Journal of Child Neurology, 2020, 14, 111-116.	0.3	1
39	Waardenburg syndrome type 2A in a large Iranian family with a novel MITF gene mutation. BMC Medical Genomics, 2021, 14, 230.	1.5	0
40	Setting up Multiplex Panels for Genetic Testing of Familial Hypertrophic Cardiomyopathy Based on Linkage Analysis. Iranian Journal of Public Health, 2016, 45, 329-39.	0.5	0
41	Next-generation sequencing identified novel truncating mutations in BBS9 causing Bardet Biedl syndrome in two Iranian consanguineous families.. Iranian Journal of Child Neurology, 2022, 16, 123-133.	0.3	0