Shadab Salehpour

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/2527651/publications.pdf

Version: 2024-02-01

27	248	1162889	996849
papers	citations	8	15
		h-index	g-index

28	28	28	409
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A Double-Blind, Placebo-Controlled Comparison of Letrozole to Oxandrolone Effects upon Growth and Puberty of Children with Constitutional Delay of Puberty and Idiopathic Short Stature. Hormone Research in Paediatrics, 2010, 74, 428-435.	0.8	51
2	Cyclic Pamidronate Therapy in Children with Osteogenesis Imperfecta. Journal of Pediatric Endocrinology and Metabolism, 2010, 23, 73-80.	0.4	29
3	Effects of Miglustat on Stabilization of Neurological Disorder in Niemannâ€“Pick Disease Type C. Journal of Child Neurology, 2013, 28, 1599-1606.	0.7	27
4	Craniosynostosis in a patient with 2q37.3 deletion 5q34 duplication: Association of extra copy of MSX2 with craniosynostosis. American Journal of Medical Genetics, Part A, 2009, 149A, 1544-1549.	0.7	24
5	Four Years of Diagnostic Challenges with Tetrahydrobiopterin Deficiencies in Iranian Patients. JIMD Reports, 2016, 32, 7-14.	0.7	16
6	Molecular genetics of a cohort of 635 cases of phenylketonuria in a consanguineous population. Journal of Inherited Metabolic Disease, 2018, 41, 1159-1167.	1.7	14
7	ELMO Domain Containing 1 (ELMOD1) Gene Mutation Is Associated with Mental Retardation and Autism Spectrum Disorder. Journal of Molecular Neuroscience, 2019, 69, 312-315.	1.1	12
8	Clinical, biochemical and molecular features of Iranian families with mucopolysaccharidosis: A case series. Clinica Chimica Acta, 2017, 474, 88-95.	0.5	10
9	Mutations in the VPS13B Gene in Iranian Patients with Different Phenotypes of Cohen Syndrome. Journal of Molecular Neuroscience, 2020, 70, 21-25.	1.1	9
10	Molecular Characterization of QDPR Gene in Iranian Families with BH4 Deficiency: Reporting Novel and Recurrent Mutations. JIMD Reports, 2015, 21, 123-128.	0.7	8
11	Co-existence of phenylketonuria either with maple syrup urine disease or Sandhoff disease in two patients from Iran: emphasizing the role of consanguinity. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 1215-1219.	0.4	7
12	Whole-Exome Sequencing Uncovers Novel Causative Variants and Additional Findings in Three Patients Affected by Glycogen Storage Disease Type VI and Fanconiâ˜Bickel Syndrome. Frontiers in Genetics, 2020, 11, 601566.	1.1	7
13	Association of a Novel Nonsense Mutation in KIAA1279 with Goldberg-Shprintzen Syndrome. Iranian Journal of Child Neurology, 2017, 11, 70-74.	0.2	7
14	Three novel mutations in Iranian patients with Tay-Sachs disease. Iranian Biomedical Journal, 2014, 18, 114-9.	0.4	5
15	Identification of a novel de novo mutation in the CTNNB1 gene in an Iranian patient with intellectual disability. Neurological Sciences, 2022, 43, 2859.	0.9	4
16	A novel splice site mutation in the GNPTAB gene in an Iranian patient with mucolipidosis II Î±/Î². Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 991-3.	0.4	3
17	Effects of selenium supplementation on serum C reactive protein level: A systematic review and meta-analysis of randomized controlled clinical trials. Obesity Medicine, 2020, 17, 100182.	0.5	3
18	The treatment and clinical follow-up outcome in Iranian patients with tetrahydrobiopterin deficiency. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 1157-1167.	0.4	3

#	Article	IF	Citations
19	Frequency of Pediatric Acute Respiratory Tract Infections in Iran; A Systematic Review. Archives of Pediatric Infectious Diseases, 2013, 1, 44-52.	0.1	3
20	The effect of omega-3 fatty acids and vitamin E co-supplementation on glycemic profile: A systematic review and meta-analysis of randomized controlled trials. Obesity Medicine, 2020, 17, 100180.	0.5	1
21	A novelSRD5A2mutation in an Iranian family with sex development disorder. Andrologia, 2021, 53, e13847.	1.0	1
22	Analysis of the HEXA, HEXB, ARSA, and SMPD1 Genes in 68 Iranian Patients. Journal of Molecular Neuroscience, 2021, , 1.	1.1	1
23	Long-Term Effect of Nephrocalcinosis on Renal Function and Body Growth Index in Children: A Retrospective Single Center Study. Iranian Journal of Pediatrics, 2018, 28, .	0.1	1
24	Molecular characterization of a large cohort of mucopolysaccharidosis patients: Iran Mucopolysaccharidosis REâ€diagnosis study (IMPRESsion). Human Mutation, 2022, , .	1.1	1
25	Neurodegeneration with brain iron accumulation 2A: Report of four independent cases. Meta Gene, 2018, 15, 87-89.	0.3	0
26	Next generation sequencing elucidated a clinically undiagnosed metabolic disorder - An Iranian family with hereditary orotic aciduria. Gene Reports, 2019, 16, 100457.	0.4	0
27	Identification of novel mutations among Iranian NPC1 patients: a bioinformatics approach to predict pathogenic mutations. Hereditas, 2022, 159, 8.	0.5	0