

Parastoo Rostami

List of Publications by Year in descending order

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Version: 2024-02-01

19
papers

137
citations

1477746

6
h-index

1281420

11
g-index

19
all docs

19
docs citations

19
times ranked

258
citing authors

#	ARTICLE	IF	CITATIONS
1	Inborn errors of metabolism underlying primary immunodeficiencies. <i>Journal of Clinical Immunology</i> , 2014, 34, 753-771.	2.0	27
2	Clinical presentation and outcome in infantile Sandhoff disease: a case series of 25 patients from Iranian neurometabolic bioregistry with five novel mutations. <i>Orphanet Journal of Rare Diseases</i> , 2018, 13, 130.	1.2	17
3	Role of vitamin D and vitamin D receptor gene polymorphisms on residual beta cell function in children with type 1 diabetes mellitus. <i>Pharmacological Reports</i> , 2019, 71, 282-288.	1.5	15
4	The First Report of Relative Incidence of Inherited White Matter Disorders in an Asian Country Based on an Iranian Bioregistry System. <i>Journal of Child Neurology</i> , 2018, 33, 255-259.	0.7	13
5	Primary creatine deficiency syndrome as a potential missed diagnosis in children with psychomotor delay and seizure: case presentation with two novel variants and literature review. <i>Acta Neurologica Belgica</i> , 2020, 120, 511-516.	0.5	12
6	Early infantile presentation of 3-methylglutaconic aciduria type 1 with a novel mutation in AUH gene: A case report and literature review. <i>Brain and Development</i> , 2017, 39, 714-716.	0.6	7
7	AGPAT2 gene mutation in a child with Berardinelli-Seip congenital lipodystrophy syndrome. <i>Annales D'Endocrinologie</i> , 2013, 74, 59-61.	0.6	6
8	Placenta derived Mesenchymal Stem Cells transplantation in Type 1 diabetes: preliminary report of phase 1 clinical trial. <i>Journal of Diabetes and Metabolic Disorders</i> , 2021, 20, 1179-1189.	0.8	6
9	Pericentric Inversion of Chromosome 9 in an Infant With Ambiguous Genitalia. <i>Acta Medica Iranica</i> , 2017, 55, 655-657.	0.8	5
10	Molecular investigation of mutations in androgen receptor and 5 α - α -reductase α 2 genes in 46,XY Disorders of Sex Development with normal testicular development. <i>Andrologia</i> , 2019, 51, e13250.	1.0	4
11	Genotype-phenotype correlation and description of two novel mutations in Iranian patients with glycogen storage disease 1b (GSD1b). <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 35.	1.2	4
12	Molecular and clinical characterization of autoimmune polyendocrinopathy-candidiasis-ectodermal dystrophy syndrome (APECED) in Iranian non-Jewish patients: report of two novel AIRE gene pathogenic variants. <i>Orphanet Journal of Rare Diseases</i> , 2022, 17, 10.	1.2	4
13	Methylmalonic acidemia with emergency hypertension. <i>Nefrologia</i> , 2016, 36, 75-76.	0.2	3
14	A Randomized Clinical Trial of Insulin Glargine and Aspart, Compared to NPH and Regular Insulin in Children with Type 1 Diabetes Mellitus. <i>Iranian Journal of Pediatrics</i> , 2014, 24, 173-8.	0.1	3
15	Neurological and Vascular Manifestations of Ethylmalonic Encephalopathy. <i>Iranian Journal of Child Neurology</i> , 2017, 11, 57-60.	0.2	3
16	Insulin edema in a child with diabetes mellitus type 1. <i>Turkish Journal of Pediatrics</i> , 2012, 54, 309-11.	0.3	3
17	Early and delayed puberty among Iranian children with obesity. <i>Minerva Endocrinology</i> , 2020, , .	0.6	2
18	A suggested screening method for hypothyroidism in very preterm and/or very low birth weight neonates. <i>Revista Paulista De Pediatria</i> , 2022, 40, e2020376.	0.4	2

#	ARTICLE	IF	CITATIONS
19	Methylmalonic acidemia with emergency hypertension. <i>Nefrologia</i> , 2016, 36, 75-76.	0.2	1