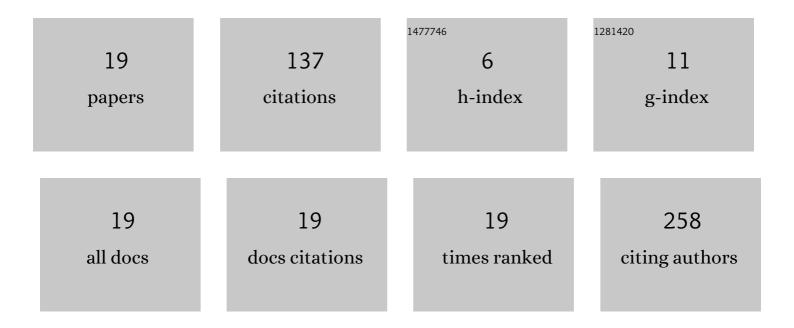
Parastoo Rostami

List of Publications by Year in descending order

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

#	Article	IF	CITATIONS
19	Insulin edema in a child with diabetes mellitus type 1. Turkish Journal of Pediatrics, 2012, 54, 309-11.	0.3	3