# Parastoo Rostami

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/2246772/publications.pdf

Version: 2024-02-01

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|---|---|---|---|
| 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 | 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 |