Shahab Noorian

List of Publications by Year in Descending Order

14 papers	24 citations	3 h-index	4 g-index
18 ext. papers	38 ext. citations	1.4 avg, IF	1.24 L-index

#	Paper	IF	Citations
14	Compound Heterozygous Mutations Presented with Quadriparesis and Menopause. A Case Report.. *Twin Research and Human Genetics*, **2022**, 1-3	2.2	
13	A novel pathogenic variant of gene causes rigidity and multifocal seizure syndrome, lethal neonatal. *International Journal of Neuroscience*, **2021**, 131, 875-878	2	3
12	Whole-Exome Sequencing in Idiopathic Short Stature: Rare Mutations Affecting Growth. *Journal of Pediatric Genetics*, **2021**, 10, 284-291	0.7	1
11	The Prevalence of Selective and Partial Immunoglobulin A Deficiency in Patients with Autoimmune Polyendocrinopathy. *Immunological Investigations*, **2021**, 1-9	2.9	1
10	A Family With Novel X-Linked Recessive Homozygous Mutation in (c.628_629 del, p.1210fs*) in Kallmann Syndrome Associated Unilateral Ptosis: Case Report and Literature Review. *AACE Clinical Case Reports*, **2021**, 7, 216-219	0.7	1
9	The Role of Thyroid Function Tests in Diagnosing Allan-herndon-dudley Syndrome Revisited: A Novel Iran-based Mutation.. *Basic and Clinical Neuroscience*, **2021**, 12, 563-568	1.4	0
8	COVID-19 and Diabetic Ketoacidosis in a Child: A Case Report. *International Journal of Enteric Pathogens*, **2021**, 9, 78-80	0.2	
7	Diagnostic Approach to the Patients with Suspected Primary Immunodeficiency. *Endocrine, Metabolic and Immune Disorders - Drug Targets*, **2020**, 20, 157-171	2.2	6
6	A novel missense mutation of the HGD gene causes Alkaptonuria. *Meta Gene*, **2018**, 18, 174-176	0.7	0
5	A novel nonsense mutation in the WFS1 gene causes the Wolfram syndrome. *Journal of Pediatric Endocrinology and Metabolism*, **2016**, 29, 607-9	1.6	2
4	Dizygotic Twins Concordant for Down Syndrome: Implication for Establishing a National Birth Defect Registry in Iran. *Iranian Journal of Public Health*, **2016**, 45, 1667-1668	0.7	
3	GCK Mutation in a Child with Maturity Onset Diabetes of the Young, Type 2. *Iranian Journal of Pediatrics*, **2013**, 23, 226-8	1	3
2	Reliability of pubertal self assessment method: an Iranian study. *Iranian Journal of Pediatrics*, **2013**, 23, 327-32	1	7
1	Familial hypercholesterolemia in an Iranian family due to a mutation in the APOE gene (first case report). *Journal of Diabetes and Metabolic Disorders*,1	2.5	