Shahab Noorian

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/8735933/publications.pdf

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16	48	1937632 4	1720014 7
papers	citations	h-index	g-index

18	18	18	116
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Diagnostic Approach to the Patients with Suspected Primary Immunodeficiency. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2020, 20, 157-171.	1.2	15
2	A novel pathogenic variant of BRAT1 gene causes rigidity and multifocal seizure syndrome, lethal neonatal. International Journal of Neuroscience, 2021, 131, 875-878.	1.6	8
3	Reliability of pubertal self assessment method: an Iranian study. Iranian Journal of Pediatrics, 2013, 23, 327-32.	0.3	8
4	GCK Mutation in a Child with Maturity Onset Diabetes of the Young, Type 2. Iranian Journal of Pediatrics, 2013, 23, 226-8.	0.3	4
5	The Prevalence of Selective and Partial Immunoglobulin A Deficiency in Patients with Autoimmune Polyendocrinopathy. Immunological Investigations, 2022, 51, 778-786.	2.0	3
6	Whole-Exome Sequencing in Idiopathic Short Stature: Rare Mutations Affecting Growth. Journal of Pediatric Genetics, 2021, 10, 284-291.	0.7	3
7	A novel nonsense mutation in the WFS1 gene causes the Wolfram syndrome. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 607-9.	0.9	2
8	A Family With Novel X-Linked Recessive Homozygous Mutation in ANOS1 (c.628_629 del, p.1210fsâˆ—) in Kallmann Syndrome Associated Unilateral Ptosis: Case Report and Literature Review. AACE Clinical Case Reports, 2021, 7, 216-219.	1.1	2
9	A novel missense mutation of the HGD gene causes Alkaptonuria. Meta Gene, 2018, 18, 174-176.	0.6	1
10	The Role of Thyroid Function Tests in the Diagnosis of Allan-Herndon-Dudley Syndrome Revisited: A Novel Mutation From Iran. Basic and Clinical Neuroscience, 2021, 12, 563-568.	0.6	1
11	Late infantile form of multiple sulfatase deficiency. Endocrinology, Diabetes and Metabolism Case Reports, 2020, 2020, .	0.5	1
12	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.	0.5	0
13	Late infantile form of multiple sulfatase deficiency. Endocrinology, Diabetes and Metabolism Case Reports, 2020, 2020, .	0.5	0
14	Familial hypercholesterolemia in an Iranian family due to a mutation in the APOE gene (first case) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 2	1.9	0
15	Compound Heterozygous Mutations Presented with Quadriparesis and Menopause. A Case Report. Twin Research and Human Genetics, 2022, , 1-3.	0.6	0
16	COVID-19 and Diabetic Ketoacidosis in a Child: A Case Report. International Journal of Enteric Pathogens, 2021, 9, 78-80.	0.1	0