Farzaneh Abbasi

List of Publications by Year
in descending order

28 papers	376 citations	759233 12 h-index	794594 19 g-index
29 all docs	29 docs citations	29 times ranked	744 citing authors

#	Article	IF	Citations
1	Insulin adherence in patients with diabetes: Risk factors for injection omission. Primary Care Diabetes, 2014, 8, 338-345.	1.8	76
2	A randomized placebo-controlled trial of alphacalcidol on the preservation of beta cell function in children with recent onset type 1 diabetes. Clinical Nutrition, 2013, 32, 911-917.	5.0	52
3	Serum IL-17, IL-23, and TGF-Î²Levels in Type 1 and Type 2 Diabetic Patients and Age-Matched Healthy Controls. BioMed Research International, 2014, 2014, 1-7.	1.9	50
4	TGF-Î² and IL-23 gene expression in unstimulated PBMCs of patients with diabetes. Endocrine, 2012, 41, 430-434.	2.3	23
5	A Genotype-First Approach for Clinical and Genetic Evaluation of Wolcott-Rallison Syndrome in a Large Cohort of Iranian Children With Neonatal Diabetes. Canadian Journal of Diabetes, 2018, 42, 272-275.	0.8	19
6	Assessment of Vitamin D Status and Response to Vitamin D3 in Obese and Non-Obese Iranian Children. Journal of Tropical Pediatrics, 2016, 62, 269-275.	1.5	18
7	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.	3.3	15
8	Vitamin D status and associated factors in recent-onset type 1 diabetic children in Iran. Journal of Diabetes and Metabolic Disorders, 2012, 11, 12.	1.9	13
9	HLA-DRB, -DQA, and DQB alleles andÂhaplotypes in Iranian patients with diabetes mellitus type I. Pediatric Diabetes, 2013, 14, 366-371.	2.9	13
10	Cardiac malformations in fetuses of gestational and pre gestational diabetic mothers. Iranian Journal of Pediatrics, 2013, 23, 664-8.	0.3	13
11	A new missense mutation in FGF23 gene in a male with hyperostosisâ€“hyperphosphatemia syndrome (HHS). Gene, 2014, 542, 269-271.	2.2	12
12	PTPN22 Single-Nucleotide Polymorphisms in Iranian Patients with Type 1 Diabetes Mellitus. Immunological Investigations, 2017, 46, 409-418.	2.0	12
13	Detection of KCNJ11 Gene Mutations in a Family with Neonatal Diabetes Mellitus. Molecular Diagnosis and Therapy, 2012, 16, 109-114.	3.8	9
14	Segregation of a novel homozygous 6 nucleotide deletion in GLUT2 gene in a Fanconiâ€“Bickel syndrome family. Gene, 2015, 557, 103-105.	2.2	9
15	Reliability of pubertal self assessment method: an Iranian study. Iranian Journal of Pediatrics, 2013, 23, 327-32.	0.3	8
16	Autoimmune polyendocrinopathy-candidiasis-ectodermal dystrophy: report of three cases from Iran. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 979-83.	0.9	5
17	Molecular investigation of WFS1 gene exon 8 in Iranian patients with Wolfram syndrome. International Journal of Diabetes in Developing Countries, 2016, 36, 75-80.	0.8	5
18	Investigating Genetic Mutations in a Large Cohort of Iranian Patients with Congenital Hyperinsulinism. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2022, 14, 87-95.	0.9	5

#	Article	IF	Citations
19	The Effect of Metformin as an Adjunct Therapy in Adolescents with Type 1 Diabetes. Journal of Clinical and Diagnostic Research JCDR, 2017, 11, SC01-SC04.	0.8	4
20	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.	2.7	4
21	Comparison of Bone Mineral Density in Common Variable Immunodeficiency and X-Linked Agammaglobulinaemia Patients. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2017, 17, 134-140.	1.2	3
22	Hyperostosis-hyperphosphatemia syndrome (HHS): report of two cases with a recurrent mutation and review of the literature. Journal of Pediatric Endocrinology and Metabolism, 2015, 28, 231-5.	0.9	2
23	Early and delayed puberty among Iranian children with obesity. Minerva Endocrinology, 2020, , .	1.1	2
24	A case of H syndrome with a novel mutation in SLC29A3. Meta Gene, 2019, 21, 100599.	0.6	1
25	p.Gln318X and p.Val281Leu as the Major Variants of CYP21A2 Gene in Children with Idiopathic Premature Pubarche. International Journal of Endocrinology, 2020, 2020, 1-9.	1.5	1
26	Proptosis, Micrognathia, Low Set Ear and Chest Deformity in a Patient with Extra Marker Chromosome 22. Acta Medica Iranica, 2015, 53, 782-4.	0.8	1
27	Indole-based derivatives effect on rats with polycystic ovary syndrome. Journal of Pharmaceutical Investigation, 2017, 47, 439-444.	5.3	0
28	Evaluation of Patients Referred to Children's Medical Center Laboratory for Diagnosis of Mucopolysaccharidoses: Eight Years' Experience from Iran. Journal of Child Science, 2021, 11, e299-e305.	0.2	0