

Chun Lin Wang

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

14
papers

51
citations

4
h-index

6
g-index

23
ext. papers

87
ext. citations

3.3
avg, IF

1.75
L-index

#	Paper	IF	Citations
14	Digenetic inheritance of SLC12A3 and CLCNKB genes in a Chinese girl with Gitelman syndrome. <i>BMC Pediatrics</i> , 2019 , 19, 114	2.6	12
13	U-shaped relationship between birth weight and childhood blood pressure in China. <i>BMC Pediatrics</i> , 2019 , 19, 264	2.6	11
12	Association between VEGF-460T/C gene polymorphism and clinical outcomes of nasopharyngeal carcinoma treated with intensity-modulated radiation therapy. <i>OncoTargets and Therapy</i> , 2017 , 10, 909-918	4.4	7
11	GCK mutations in Chinese MODY2 patients: a family pedigree report and review of Chinese literature. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2016 , 29, 959-64	1.6	5
10	Regional Disparities in Obesity Among a Heterogeneous Population of Chinese Children and Adolescents. <i>JAMA Network Open</i> , 2021 , 4, e2131040	10.4	4
9	Associations between maternal age at menarche and anthropometric and metabolic parameters in the adolescent offspring. <i>Clinical Endocrinology</i> , 2019 , 90, 702-710	3.4	3
8	A Novel Compound Heterozygous Variant Causes 17 β -Hydroxylase/17, 20-Lyase Deficiency. <i>Frontiers in Genetics</i> , 2019 , 10, 996	4.5	3
7	Association of Vitamin D Receptor Gene Polymorphisms with Metabolic Syndrome in Chinese Children. <i>International Journal of General Medicine</i> , 2021 , 14, 57-66	2.3	3
6	Neonatal presentation of familial glucocorticoid deficiency with a MRAP mutation: A case report. <i>Molecular Genetics and Metabolism Reports</i> , 2016 , 9, 15-7	1.8	2
5	The first central precocious puberty proteomic profiles revealed multiple metabolic networks and novel key disease-associated proteins. <i>Aging</i> , 2021 , 13, 24236-24250	5.6	0
4	High prevalence of elevated serum liver enzymes in Chinese children suggests metabolic syndrome as a common risk factor. <i>Journal of Paediatrics and Child Health</i> , 2020 , 56, 1590-1596	1.3	0
3	Premature ovarian insufficiency in children: Etiology, clinical management and treatment. <i>Journal of Advanced Pediatrics and Child Health</i> , 2020 , 3, 047-055	0	0
2	A 15q25.2 microdeletion phenotype for premature ovarian failure in a Chinese girl: a case report and review of literature. <i>BMC Medical Genomics</i> , 2020 , 13, 126	3.7	0
1	A Novel de Novo Variant in 5' UTR of the NIPBL Associated with Cornelia de Lange Syndrome. <i>Genes</i> , 2022 , 13, 740	4.2	0