Chun Lin Wang

List of Publications by Year in Descending Order

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	A Novel de Novo Variant in 5? UTR of the NIPBL Associated with Cornelia de Lange Syndrome. *Genes*, **2022**, 13, 740	4.2	
13	The first central precocious puberty proteomic profiles revealed multiple metabolic networks and novel key disease-associated proteins. *Aging*, **2021**, 13, 24236-24250	5.6	0
12	Regional Disparities in Obesity Among a Heterogeneous Population of Chinese Children and Adolescents. *JAMA Network Open*, **2021**, 4, e2131040	10.4	4
11	Association of Vitamin D Receptor Gene Polymorphisms with Metabolic Syndrome in Chinese Children. *International Journal of General Medicine*, **2021**, 14, 57-66	2.3	3
10	Premature ovarian insufficiency in children: Etiology, clinical management and treatment. *Journal of Advanced Pediatrics and Child Health*, **2020**, 3, 047-055	0	
9	High prevalence of elevated serum liver enzymes in Chinese children suggests metabolic syndrome as a common risk factor. *Journal of Paediatrics and Child Health*, **2020**, 56, 1590-1596	1.3	0
8	A 15q25.2 microdeletion phenotype for premature ovarian failure in a Chinese girl: a case report and review of literature. *BMC Medical Genomics*, **2020**, 13, 126	3.7	
7	Digenetic inheritance of SLC12A3 and CLCNKB genes in a Chinese girl with Gitelman syndrome. *BMC Pediatrics*, **2019**, 19, 114	2.6	12
6	Associations between maternal age at menarche and anthropometric and metabolic parameters in the adolescent offspring. *Clinical Endocrinology*, **2019**, 90, 702-710	3.4	3
5	U-shaped relationship between birth weight and childhood blood pressure in China. *BMC Pediatrics*, **2019**, 19, 264	2.6	11
4	A Novel Compound Heterozygous Variant Causes 17-Hydroxylase/17, 20-Lyase Deficiency. *Frontiers in Genetics*, **2019**, 10, 996	4.5	3
3	Association between VEGF-460T/C gene polymorphism and clinical outcomes of nasopharyngeal carcinoma treated with intensity-modulated radiation therapy. *OncoTargets and Therapy*, **2017**, 10, 909-918	4.4	7
2	GCK mutations in Chinese MODY2 patients: a family pedigree report and review of Chinese literature. *Journal of Pediatric Endocrinology and Metabolism*, **2016**, 29, 959-64	1.6	5
1	Neonatal presentation of familial glucocorticoid deficiency with a MRAP mutation: A case report. *Molecular Genetics and Metabolism Reports*, **2016**, 9, 15-7	1.8	2