

# Chun Gong

## List of Publications by Year in descending order

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Version: 2024-02-01

52  
papers

494  
citations

777949

13  
h-index

939365

18  
g-index

72  
all docs

72  
docs citations

72  
times ranked

790  
citing authors

#	ARTICLE	IF	CITATIONS
1	Genetic and Phenotypic Spectrum of KBC Syndrome: A Report of 13 New Chinese Cases and a Review of the Literature. <i>Journal of Personalized Medicine</i> , 2022, 12, 407.	1.1	7
2	Pediatric Continuous Reference Intervals of Serum Insulin-like Growth Factor 1 Levels in a Healthy Chinese Children Population “ Based on PRINCE Study. <i>Endocrine Practice</i> , 2022, 28, 696-702.	1.1	8
3	First Clinical Study on Long-Acting Growth Hormone Therapy in Children with Turner Syndrome. <i>Hormone and Metabolic Research</i> , 2022, 54, 389-395.	0.7	2
4	Growth Curves of Chinese Children with Androgen Insensitivity Syndrome: A Multicenter Registry Study. <i>Journal of Personalized Medicine</i> , 2022, 12, 771.	1.1	0
5	DNA Hypermethylation and a Specific Methylation Spectrum on the X Chromosome in Turner Syndrome as Determined by Nanopore Sequencing. <i>Journal of Personalized Medicine</i> , 2022, 12, 872.	1.1	0
6	Correlation Analysis of Genotypes and Phenotypes in Chinese Male Pediatric Patients With Congenital Hypogonadotropic Hypogonadism. <i>Frontiers in Endocrinology</i> , 2022, 13, .	1.5	2
7	Whole exome sequencing for non-selective pediatric patients with hyperlipidemia. <i>Gene</i> , 2021, 768, 145310.	1.0	2
8	Clinical and genetic characteristics of hypophosphatasia in Chinese children. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 159.	1.2	5
9	Heterozygous Recurrent Mutations Inducing Dysfunction of ROR2 Gene in Patients With Short Stature. <i>Frontiers in Cell and Developmental Biology</i> , 2021, 9, 661747.	1.8	4
10	Exome sequencing reveals genetic architecture in patients with isolated or syndromic short stature. <i>Journal of Genetics and Genomics</i> , 2021, 48, 396-402.	1.7	21
11	High Prevalence of Obesity but Low Physical Activity in Children Aged 9“11 Years in Beijing. <i>Diabetes, Metabolic Syndrome and Obesity: Targets and Therapy</i> , 2021, Volume 14, 3323-3335.	1.1	6
12	Whole Exome Sequencing Uncovered the Genetic Architecture of Growth Hormone Deficiency Patients. <i>Frontiers in Endocrinology</i> , 2021, 12, 711991.	1.5	6
13	Associations Between Sleep Duration, Wake-Up Time, Bedtime, and Abdominal Obesity: Results From 9559 Chinese Children Aged 7“18 Years. <i>Frontiers in Endocrinology</i> , 2021, 12, 735952.	1.5	7
14	One hundred twelve cases of 46, XY DSD patients after initial gender assignment: a short-term survey of gender role and gender dysphoria. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 416.	1.2	3
15	Hormonal changes throughout puberty in boys. <i>Chinese Medical Journal</i> , 2021, Publish Ahead of Print, 362-364.	0.9	0
16	SOX2 heterozygous mutations cause multiple extraocular phenotypes in boys. <i>Chinese Medical Journal</i> , 2021, Publish Ahead of Print, .	0.9	1
17	Editorial: Childhood Diabetes in Low- and Middle-Income Countries. <i>Frontiers in Endocrinology</i> , 2021, 12, 830700.	1.5	0
18	Clinical, Biochemical, Molecular, and Outcome Features of Mitochondrial 3-Hydroxy-3-Methylglutaryl-CoA Synthase Deficiency in 10 Chinese Patients. <i>Frontiers in Genetics</i> , 2021, 12, 816779.	1.1	2

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19	Exploring the efficacy of testosterone undecanoate in male children with 5 $\alpha$ -reductase deficiency. <i>Pediatric Investigation</i> , 2021, 5, 249-254.	0.6	3
20	A novel heterozygous MKRN3 nonsense mutation in a Chinese girl with idiopathic central precocious puberty. <i>Medicine (United States)</i> , 2020, 99, e22295.	0.4	5
21	The first familial NSD2 cases with a novel variant in a Chinese father and daughter with atypical WHS facial features and a 7.5-year follow-up of growth hormone therapy. <i>BMC Medical Genomics</i> , 2020, 13, 181.	0.7	6
22	Clinical characteristics and genotype-phenotype correlations of 130 Chinese children in a high-homogeneity single-center cohort with 5 $\alpha$ -reductase 2 deficiency. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2020, 8, e1431.	0.6	17
23	Aromatase deficiency: A case series of 46, XX Chinese children and a systematic review of the literature. <i>Clinical Endocrinology</i> , 2020, 93, 687-695.	1.2	3
24	Central precocious puberty as a prelude to hypogonadism in a patient with Klinefelter syndrome. <i>Pediatric Investigation</i> , 2019, 3, 127-130.	0.6	3
25	An Evaluation of the Accuracy of a Flash Glucose Monitoring System in Children with Diabetes in comparison with Venous Blood Glucose. <i>Journal of Diabetes Research</i> , 2019, 2019, 1-7.	1.0	9
26	New insights into 5 $\alpha$ -reductase type 2 deficiency based on a multi-centre study: regional distribution and genotype-phenotype profiling of <i>SRD5A2</i> in 190 Chinese patients. <i>Journal of Medical Genetics</i> , 2019, 56, 685-692.	1.5	16
27	Growth Pattern in Chinese Children With 5 $\alpha$ -Reductase Type 2 Deficiency: A Retrospective Multicenter Study. <i>Frontiers in Pharmacology</i> , 2019, 10, 173.	1.6	4
28	Variant analysis of the chromodomain helicase DNA-binding protein 7 in pediatric disorders of sex development. <i>Pediatric Investigation</i> , 2019, 3, 31-38.	0.6	1
29	A cross-sectional survey of adrenal steroid hormones among overweight/obese boys according to puberty stage. <i>BMC Pediatrics</i> , 2019, 19, 414.	0.7	7
30	Kabuki syndrome: novel pathogenic variants, new phenotypes and review of literature. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 255.	1.2	24
31	Novel phenotypes and genotypes in Antley-Bixler syndrome caused by cytochrome P450 oxidoreductase deficiency: based on the first cohort of Chinese children. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 299.	1.2	12
32	SUN-LB041 Clinical and Molecular Characteristics, Genotype-Phenotype Correlation in 113 Chinese Children with <i>SRD5A2</i> Gene Mutations. <i>Journal of the Endocrine Society</i> , 2019, 3, .	0.1	0
33	Novel pathogenic <i>RECQL4</i> variants in Chinese patients with Rothmund-Thomson syndrome. <i>Gene</i> , 2018, 654, 110-115.	1.0	8
34	Phenotype and Molecular Characterizations of 30 Children From China With <i>NR5A1</i> Mutations. <i>Frontiers in Pharmacology</i> , 2018, 9, 1224.	1.6	14
35	Validity of web-based self-assessment of pubertal development against pediatrician assessments. <i>Pediatric Investigation</i> , 2018, 2, 141-148.	0.6	4
36	Clinical and Molecular Spectrum of Glutamate Dehydrogenase Gene Defects in 26 Chinese Congenital Hyperinsulinemia Patients. <i>Journal of Diabetes Research</i> , 2018, 2018, 1-6.	1.0	10

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37	Type 1 diabetes mellitus care and education in China: The 3C study of coverage, cost, and care in Beijing and Shantou. <i>Diabetes Research and Clinical Practice</i> , 2017, 129, 32-42.	1.1	27
38	Genome-wide analysis of differential DNA methylation in Silver-Russell syndrome. <i>Science China Life Sciences</i> , 2017, 60, 692-699.	2.3	17
39	Prenatal and early diagnosis of Chinese 3-M syndrome patients with novel pathogenic variants. <i>Clinica Chimica Acta</i> , 2017, 474, 159-164.	0.5	9
40	Permanent neonatal diabetes caused by abnormalities in chromosome 6q24. <i>Diabetic Medicine</i> , 2017, 34, 1800-1804.	1.2	8
41	Clinical and genetic features of 64 young male paediatric patients with congenital hypogonadotropic hypogonadism. <i>Clinical Endocrinology</i> , 2017, 87, 757-766.	1.2	31
42	Persistent Müllerian duct syndrome: A case report and review. <i>Experimental and Therapeutic Medicine</i> , 2017, 14, 5779-5784.	0.8	16
43	Fulminant Type 1 Diabetes in Children: A Multicenter Study in China. <i>Journal of Diabetes Research</i> , 2017, 2017, 1-6.	1.0	6
44	Increase in Peripheral Blood Intermediate Monocytes is Associated with the Development of Recent-Onset Type 1 Diabetes Mellitus in Children. <i>International Journal of Biological Sciences</i> , 2017, 13, 209-218.	2.6	25
45	Comorbidity in Adult Patients Hospitalized with Type 2 Diabetes in Northeast China: An Analysis of Hospital Discharge Data from 2002 to 2013. <i>BioMed Research International</i> , 2016, 2016, 1-9.	0.9	14
46	Genetic Analysis and Follow-Up of 25 Neonatal Diabetes Mellitus Patients in China. <i>Journal of Diabetes Research</i> , 2016, 2016, 1-9.	1.0	28
47	A randomised, open-label study of insulin glargine or neutral protamine Hagedorn insulin in Chinese paediatric patients with type 1 diabetes mellitus. <i>BMC Endocrine Disorders</i> , 2016, 16, 67.	0.9	6
48	Identification of AAAS gene mutation in Allgrove syndrome: A report of three cases. <i>Experimental and Therapeutic Medicine</i> , 2015, 10, 1277-1282.	0.8	16
49	Trends in Childhood Type 1 Diabetes Mellitus Incidence in Beijing from 1995 to 2010: A Retrospective Multicenter Study Based on Hospitalization Data. <i>Diabetes Technology and Therapeutics</i> , 2015, 17, 159-165.	2.4	32
50	Two case reports of severe pediatric hyperosmolar hyperglycemia and diabetic ketoacidosis accompanied with rhabdomyolysis and acute renal failure. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2014, 27, 1227-31.	0.4	6
51	Autologous hematopoietic stem cell transplantation and conventional insulin therapy in the treatment of children with newly diagnosed type 1 diabetes: long term follow-up. <i>Chinese Medical Journal</i> , 2014, 127, 2618-22.	0.9	9
52	Trends in the Incidence of Childhood Type 1 Diabetes Mellitus in Beijing Based on Hospitalization Data from 1995 to 2010. <i>Hormone Research in Paediatrics</i> , 2013, 80, 328-334.	0.8	13