Chun Gong

List of Publications by Year
in descending order

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Version: 2024-02-01

52	494	687363	839539
papers	citations	13	18
		h-index	g-index

72	72	72	763
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Trends in Childhood Type 1 Diabetes Mellitus Incidence in Beijing from 1995 to 2010: A Retrospective Multicenter Study Based on Hospitalization Data. Diabetes Technology and Therapeutics, 2015, 17, 159-165.	4.4	32
2	Clinical and genetic features of 64 young male paediatric patients with congenital hypogonadotropic hypogonadism. Clinical Endocrinology, 2017, 87, 757-766.	2.4	31
3	Genetic Analysis and Follow-Up of 25 Neonatal Diabetes Mellitus Patients in China. Journal of Diabetes Research, 2016, 2016, 1-9.	2.3	28
4	Type 1 diabetes mellitus care and education in China: The 3C study of coverage, cost, and care in Beijing and Shantou. Diabetes Research and Clinical Practice, 2017, 129, 32-42.	2.8	27
5	Increase in Peripheral Blood Intermediate Monocytes is Associated with the Development of Recent-Onset Type 1 Diabetes Mellitus in Children. International Journal of Biological Sciences, 2017, 13, 209-218.	6.4	25
6	Kabuki syndrome: novel pathogenic variants, new phenotypes and review of literature. Orphanet Journal of Rare Diseases, 2019, 14, 255.	2.7	24
7	Exome sequencing reveals genetic architecture in patients with isolated or syndromic short stature. Journal of Genetics and Genomics, 2021, 48, 396-402.	3.9	21
8	Genome-wide analysis of differential DNA methylation in Silver-Russell syndrome. Science China Life Sciences, 2017, 60, 692-699.	4.9	17
9	Clinical characteristics and genotypeâ€phenotype correlations of 130 Chinese children in a highâ€homogeneity singleâ€center cohort with 5Î±â€reductase 2 deficiency. Molecular Genetics & Genomic Medicine, 2020, 8, e1431.	1.2	17
10	Identification of AAAS gene mutation in Allgrove syndrome: A report of three cases. Experimental and Therapeutic Medicine, 2015, 10, 1277-1282.	1.8	16
11	Persistent MÃ¯Â¿Â½llerian duct syndrome: A case report and review. Experimental and Therapeutic Medicine, 2017, 14, 5779-5784.	1.8	16
12	New insights into 5Î±-reductase type 2 deficiency based on a multi-centre study: regional distribution and genotypeâ€“phenotype profiling ofSRD5A2in 190 Chinese patients. Journal of Medical Genetics, 2019, 56, 685-692.	3.2	16
13	Comorbidity in Adult Patients Hospitalized with Type 2 Diabetes in Northeast China: An Analysis of Hospital Discharge Data from 2002 to 2013. BioMed Research International, 2016, 2016, 1-9.	1.9	14
14	Phenotype and Molecular Characterizations of 30 Children From China With NR5A1 Mutations. Frontiers in Pharmacology, 2018, 9, 1224.	3.5	14
15	Trends in the Incidence of Childhood Type 1 Diabetes Mellitus in Beijing Based on Hospitalization Data from 1995 to 2010. Hormone Research in Paediatrics, 2013, 80, 328-334.	1.8	13
16	Novel phenotypes and genotypes in Antley-Bixler syndrome caused by cytochrome P450 oxidoreductase deficiency: based on the first cohort of Chinese children. Orphanet Journal of Rare Diseases, 2019, 14, 299.	2.7	12
17	Clinical and Molecular Spectrum of Glutamate Dehydrogenase Gene Defects in 26 Chinese Congenital Hyperinsulinemia Patients. Journal of Diabetes Research, 2018, 2018, 1-6.	2.3	10
18	Prenatal and early diagnosis of Chinese 3-M syndrome patients with novel pathogenic variants. Clinica Chimica Acta, 2017, 474, 159-164.	1.1	9

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19	An Evaluation of the Accuracy of a Flash Glucose Monitoring System in Children with Diabetes in comparison with Venous Blood Glucose. Journal of Diabetes Research, 2019, 2019, 1-7.	2.3	9
20	Autologous hematopoietic stem cell transplantation and conventional insulin therapy in the treatment of children with newly diagnosed type 1 diabetes: long term follow-up. Chinese Medical Journal, 2014, 127, 2618-22.	2.3	9
21	Permanent neonatal diabetes caused by abnormalities in chromosome 6q24. Diabetic Medicine, 2017, 34, 1800-1804.	2.3	8
22	Novel pathogenic RECQL4 variants in Chinese patients with Rothmund-Thomson syndrome. Gene, 2018, 654, 110-115.	2.2	8
23	Pediatric Continuous Reference Intervals of Serum Insulin-like Growth Factor 1 Levels in a Healthy Chinese Children Population â€“ Based on PRINCE Study. Endocrine Practice, 2022, 28, 696-702.	2.1	8
24	A cross-sectional survey of adrenal steroid hormones among overweight/obese boys according to puberty stage. BMC Pediatrics, 2019, 19, 414.	1.7	7
25	Associations Between Sleep Duration, Wake-Up Time, Bedtime, and Abdominal Obesity: Results From 9559 Chinese Children Aged 7â€“18 Years. Frontiers in Endocrinology, 2021, 12, 735952.	3.5	7
26	Genetic and Phenotypic Spectrum of KBG Syndrome: A Report of 13 New Chinese Cases and a Review of the Literature. Journal of Personalized Medicine, 2022, 12, 407.	2.5	7
27	Two case reports of severe pediatric hyperosmolar hyperglycemia and diabetic ketoacidosis accompanied with rhabdomyolysis and acute renal failure. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 1227-31.	0.9	6
28	A randomised, open-labelstudy of insulin glargine or neutral protamine Hagedorn insulin in Chinese paediatric patients with type 1 diabetes mellitus. BMC Endocrine Disorders, 2016, 16, 67.	2.2	6
29	Fulminant Type 1 Diabetes in Children: A Multicenter Study in China. Journal of Diabetes Research, 2017, 2017, 1-6.	2.3	6
30	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. BMC Medical Genomics, 2020, 13, 181.	1.5	6
31	High Prevalence of Obesity but Low Physical Activity in Children Aged 9â€“11 Years in Beijing. Diabetes, Metabolic Syndrome and Obesity: Targets and Therapy, 2021, Volume 14, 3323-3335.	2.4	6
32	Whole Exome Sequencing Uncovered the Genetic Architecture of Growth Hormone Deficiency Patients. Frontiers in Endocrinology, 2021, 12, 711991.	3.5	6
33	A novel heterozygous MKRN3 nonsense mutation in a Chinese girl with idiopathic central precocious puberty. Medicine (United States), 2020, 99, e22295.	1.0	5
34	Clinical and genetic characteristics of hypophosphatasia in Chinese children. Orphanet Journal of Rare Diseases, 2021, 16, 159.	2.7	5
35	Validity of webâ€based selfâ€assessment of pubertal development against pediatrician assessments. Pediatric Investigation, 2018, 2, 141-148.	1.4	4
36	Growth Pattern in Chinese Children With 5Î±-Reductase Type 2 Deficiency: A Retrospective Multicenter Study. Frontiers in Pharmacology, 2019, 10, 173.	3.5	4

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37	Heterozygous Recurrent Mutations Inducing Dysfunction of ROR2 Gene in Patients With Short Stature. Frontiers in Cell and Developmental Biology, 2021, 9, 661747.	3.7	4
38	Central precocious puberty as a prelude to hypogonadism in a patient with Klinefelter syndrome. Pediatric Investigation, 2019, 3, 127-130.	1.4	3
39	Aromatase deficiency: A case series of 46, XX Chinese children and a systematic review of the literature. Clinical Endocrinology, 2020, 93, 687-695.	2.4	3
40	One hundred twelve cases of 46, XY DSD patients after initial gender assignment: a short-term survey of gender role and gender dysphoria. Orphanet Journal of Rare Diseases, 2021, 16, 416.	2.7	3
41	Exploring the efficacy of testosterone undecanoate in male children with 5Î±â€reductase deficiency. Pediatric Investigation, 2021, 5, 249-254.	1.4	3
42	Whole exome sequencing for non-selective pediatric patients with hyperlipidemia. Gene, 2021, 768, 145310.	2.2	2
43	Clinical, Biochemical, Molecular, and Outcome Features of Mitochondrial 3-Hydroxy-3-Methylglutaryl-CoA Synthase Deficiency in 10 Chinese Patients. Frontiers in Genetics, 2021, 12, 816779.	2.3	2
44	First Clinical Study on Long-Acting Growth Hormone Therapy in Children with Turner Sydrome. Hormone and Metabolic Research, 2022, 54, 389-395.	1.5	2
45	Correlation Analysis of Genotypes and Phenotypes in Chinese Male Pediatric Patients With Congenital Hypogonadotropic Hypogonadism. Frontiers in Endocrinology, 2022, 13, .	3.5	2
46	Variant analysis of the chromodomain helicase <scp>DNA</scp>â€binding protein 7 in pediatric disorders of sex development. Pediatric Investigation, 2019, 3, 31-38.	1.4	1
47	SOX2 heterozygous mutations cause multiple extraocular phenotypes in boys. Chinese Medical Journal, 2021, Publish Ahead of Print, .	2.3	1
48	Hormonal changes throughout puberty in boys. Chinese Medical Journal, 2021, Publish Ahead of Print, 362-364.	2.3	0
49	SUN-LB041 Clinical and Molecular Characteristics, Genotype-Phenotype Correlation in 113 Chinese Children with SRD5A2 Gene Mutations. Journal of the Endocrine Society, 2019, 3, .	0.2	0
50	Editorial: Childhood Diabetes in Low- and Middle-Income Countries. Frontiers in Endocrinology, 2021, 12, 830700.	3.5	0
51	Growth Curves of Chinese Children with Androgen Insensitivity Syndrome: A Multicenter Registry Study. Journal of Personalized Medicine, 2022, 12, 771.	2.5	0
52	DNA Hypermethylation and a Specific Methylation Spectrum on the X Chromosome in Turner Syndrome as Determined by Nanopore Sequencing. Journal of Personalized Medicine, 2022, 12, 872.	2.5	0