Chun Gong

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/2999771/publications.pdf

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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	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
2	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
3	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
4	Growth Curves of Chinese Children with Androgen Insensitivity Syndrome: A Multicenter Registry Study. Journal of Personalized Medicine, 2022, 12, 771.	2.5	0
5	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
6	Correlation Analysis of Genotypes and Phenotypes in Chinese Male Pediatric Patients With Congenital Hypogonadotropic Hypogonadism. Frontiers in Endocrinology, 2022, 13, .	3.5	2
7	Whole exome sequencing for non-selective pediatric patients with hyperlipidemia. Gene, 2021, 768, 145310.	2.2	2
8	Clinical and genetic characteristics of hypophosphatasia in Chinese children. Orphanet Journal of Rare Diseases, 2021, 16, 159.	2.7	5
9	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
10	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
11	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
12	Whole Exome Sequencing Uncovered the Genetic Architecture of Growth Hormone Deficiency Patients. Frontiers in Endocrinology, 2021, 12, 711991.	3.5	6
13	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
14	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
15	Hormonal changes throughout puberty in boys. Chinese Medical Journal, 2021, Publish Ahead of Print, 362-364.	2.3	0
16	SOX2 heterozygous mutations cause multiple extraocular phenotypes in boys. Chinese Medical Journal, 2021, Publish Ahead of Print, .	2.3	1
17	Editorial: Childhood Diabetes in Low- and Middle-Income Countries. Frontiers in Endocrinology, 2021, 12, 830700.	3.5	0
18	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

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19	Exploring the efficacy of testosterone undecanoate in male children with 5Î±â€reductase deficiency. Pediatric Investigation, 2021, 5, 249-254.	1.4	3
20	A novel heterozygous MKRN3 nonsense mutation in a Chinese girl with idiopathic central precocious puberty. Medicine (United States), 2020, 99, e22295.	1.0	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. BMC Medical Genomics, 2020, 13, 181.	1.5	6
22	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
23	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
24	Central precocious puberty as a prelude to hypogonadism in a patient with Klinefelter syndrome. Pediatric Investigation, 2019, 3, 127-130.	1.4	3
25	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
26	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
27	Growth Pattern in Chinese Children With 5Î±-Reductase Type 2 Deficiency: A Retrospective Multicenter Study. Frontiers in Pharmacology, 2019, 10, 173.	3.5	4
28	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
29	A cross-sectional survey of adrenal steroid hormones among overweight/obese boys according to puberty stage. BMC Pediatrics, 2019, 19, 414.	1.7	7
30	Kabuki syndrome: novel pathogenic variants, new phenotypes and review of literature. Orphanet Journal of Rare Diseases, 2019, 14, 255.	2.7	24
31	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
32	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
33	Novel pathogenic RECQL4 variants in Chinese patients with Rothmund-Thomson syndrome. Gene, 2018, 654, 110-115.	2.2	8
34	Phenotype and Molecular Characterizations of 30 Children From China With NR5A1 Mutations. Frontiers in Pharmacology, 2018, 9, 1224.	3.5	14
35	Validity of webâ€based selfâ€assessment of pubertal development against pediatrician assessments. Pediatric Investigation, 2018, 2, 141-148.	1.4	4
36	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

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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. Diabetes Research and Clinical Practice, 2017, 129, 32-42.	2.8	27
38	Genome-wide analysis of differential DNA methylation in Silver-Russell syndrome. Science China Life Sciences, 2017, 60, 692-699.	4.9	17
39	Prenatal and early diagnosis of Chinese 3-M syndrome patients with novel pathogenic variants. Clinica Chimica Acta, 2017, 474, 159-164.	1.1	9
40	Permanent neonatal diabetes caused by abnormalities in chromosome 6q24. Diabetic Medicine, 2017, 34, 1800-1804.	2.3	8
41	Clinical and genetic features of 64 young male paediatric patients with congenital hypogonadotropic hypogonadism. Clinical Endocrinology, 2017, 87, 757-766.	2.4	31
42	Persistent MÃ¯Â¿Â½llerian duct syndrome: A case report and review. Experimental and Therapeutic Medicine, 2017, 14, 5779-5784.	1.8	16
43	Fulminant Type 1 Diabetes in Children: A Multicenter Study in China. Journal of Diabetes Research, 2017, 2017, 1-6.	2.3	6
44	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
45	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
46	Genetic Analysis and Follow-Up of 25 Neonatal Diabetes Mellitus Patients in China. Journal of Diabetes Research, 2016, 2016, 1-9.	2.3	28
47	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
48	Identification of AAAS gene mutation in Allgrove syndrome: A report of three cases. Experimental and Therapeutic Medicine, 2015, 10, 1277-1282.	1.8	16
49	Trends in Childhood Type 1 Diabetes Mellitus Incidence in Beijing from 1995 to 2010: A Retrospective Multicenter Study Based on Hospitalization Data. Diabetes Technology and Therapeutics, 2015, 17, 159-165.	4.4	32
50	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
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. Chinese Medical Journal, 2014, 127, 2618-22.	2.3	9
52	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