Chang Su

List of Publications by Year
in descending order

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20 papers	217 citations	1039406 9 h-index	1058022 14 g-index
20 all docs	20 docs citations	20 times ranked	420 citing authors

#	Article	IF	Citations
1	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.	1.1	0
2	Clinical and genetic characteristics of hypophosphatasia in Chinese children. Orphanet Journal of Rare Diseases, 2021, 16, 159.	1.2	5
3	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.	1.1	6
4	Whole Exome Sequencing Uncovered the Genetic Architecture of Growth Hormone Deficiency Patients. Frontiers in Endocrinology, 2021, 12, 711991.	1.5	6
5	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.	1.5	7
6	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.	1.0	9
7	The diagnosis of cystinosis in patients reveals new CTNS gene mutations in the Chinese population. Journal of Pediatric Endocrinology and Metabolism, 2019, 32, 375-382.	0.4	3
8	A cross-sectional survey of adrenal steroid hormones among overweight/obese boys according to puberty stage. BMC Pediatrics, 2019, 19, 414.	0.7	7
9	Kabuki syndrome: novel pathogenic variants, new phenotypes and review of literature. Orphanet Journal of Rare Diseases, 2019, 14, 255.	1.2	24
10	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.	1.2	12
11	Liquid Chromatography-Tandem Mass Spectrometry-Based Characterization of Steroid Hormone Profiles in Healthy 6 to 14-Year-Old Male Children. Chinese Medical Journal, 2018, 131, 862-866.	0.9	3
12	Pseudohypoaldosteronism Type II Caused by CUL3 Mutation Presented with Visual Impairment. Chinese Medical Journal, 2018, 131, 1879-1881.	0.9	2
13	Clinical and Molecular Spectrum of Glutamate Dehydrogenase Gene Defects in 26 Chinese Congenital Hyperinsulinemia Patients. Journal of Diabetes Research, 2018, 2018, 1-6.	1.0	10
14	A severe case of hyperinsulinism due to hemizygous activating mutation of glutamate dehydrogenase. Pediatric Diabetes, 2017, 18, 911-916.	1.2	15
15	Genome-wide analysis of differential DNA methylation in Silver-Russell syndrome. Science China Life Sciences, 2017, 60, 692-699.	2.3	17
16	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.	2.6	25
17	Genetic Analysis and Follow-Up of 25 Neonatal Diabetes Mellitus Patients in China. Journal of Diabetes Research, 2016, 2016, 1-9.	1.0	28
18	Congenital hyperinsulinism in Chinese patients: 5â€yr treatment outcome of 95 clinical cases with genetic analysis of 55 cases. Pediatric Diabetes, 2016, 17, 227-234.	1.2	19

#	Article	IF	Citations
19	Long-Term Follow-Up and Mutation Analysis of 27 Chinese Cases of Congenital Hyperinsulinism. Hormone Research in Paediatrics, 2014, 81, 169-176.	0.8	10
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.	0.9	9