

Chang Su

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7796263/publications.pdf>

Version: 2024-02-01

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	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
2	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
3	Kabuki syndrome: novel pathogenic variants, new phenotypes and review of literature. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 255.	1.2	24
4	Congenital hyperinsulinism in Chinese patients: 5-yr treatment outcome of 95 clinical cases with genetic analysis of 55 cases. <i>Pediatric Diabetes</i> , 2016, 17, 227-234.	1.2	19
5	Genome-wide analysis of differential DNA methylation in Silver-Russell syndrome. <i>Science China Life Sciences</i> , 2017, 60, 692-699.	2.3	17
6	A severe case of hyperinsulinism due to hemizygous activating mutation of glutamate dehydrogenase. <i>Pediatric Diabetes</i> , 2017, 18, 911-916.	1.2	15
7	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
8	Long-Term Follow-Up and Mutation Analysis of 27 Chinese Cases of Congenital Hyperinsulinism. <i>Hormone Research in Paediatrics</i> , 2014, 81, 169-176.	0.8	10
9	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
10	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
11	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
12	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
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	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
15	Whole Exome Sequencing Uncovered the Genetic Architecture of Growth Hormone Deficiency Patients. <i>Frontiers in Endocrinology</i> , 2021, 12, 711991.	1.5	6
16	Clinical and genetic characteristics of hypophosphatasia in Chinese children. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 159.	1.2	5
17	Liquid Chromatography-Tandem Mass Spectrometry-Based Characterization of Steroid Hormone Profiles in Healthy 6 to 14-Year-Old Male Children. <i>Chinese Medical Journal</i> , 2018, 131, 862-866.	0.9	3
18	The diagnosis of cystinosis in patients reveals new CTNS gene mutations in the Chinese population. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2019, 32, 375-382.	0.4	3

#	ARTICLE	IF	CITATIONS
19	Pseudohypoaldosteronism Type II Caused by CUL3 Mutation Presented with Visual Impairment. Chinese Medical Journal, 2018, 131, 1879-1881.	0.9	2
20	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