Lijun Fan

List of Publications by Year
in descending order

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13 papers	95 citations	1684188 5 h-index	1474206 9 g-index
16 all docs	16 docs citations	16 times ranked	76 citing authors

#	Article	IF	Citations
1	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
2	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
3	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
4	Phenotype and Molecular Characterizations of 30 Children From China With NR5A1 Mutations. Frontiers in Pharmacology, 2018, 9, 1224.	3.5	14
5	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
6	Whole Exome Sequencing Uncovered the Genetic Architecture of Growth Hormone Deficiency Patients. Frontiers in Endocrinology, 2021, 12, 711991.	3.5	6
7	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
8	Exploring the efficacy of testosterone undecanoate in male children with 5Î±â€reductase deficiency. Pediatric Investigation, 2021, 5, 249-254.	1.4	3
9	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
10	SOX2 heterozygous mutations cause multiple extraocular phenotypes in boys. Chinese Medical Journal, 2021, Publish Ahead of Print, .	2.3	1
11	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
12	Growth Curves of Chinese Children with Androgen Insensitivity Syndrome: A Multicenter Registry Study. Journal of Personalized Medicine, 2022, 12, 771.	2.5	0
13	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