

# Lijun Fan

## List of Publications by Year in descending order

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13  
papers

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citations

1683354

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1473754

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#	ARTICLE	IF	CITATIONS
1	Exome sequencing reveals genetic architecture in patients with isolated or syndromic short stature. <i>Journal of Genetics and Genomics</i> , 2021, 48, 396-402.	1.7	21
2	Clinical characteristics and genotype-phenotype correlations of 130 Chinese children in a high-homogeneity single-center cohort with 5 $\alpha$ -reductase 2 deficiency. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2020, 8, e1431.	0.6	17
3	New insights into 5 $\alpha$ -reductase type 2 deficiency based on a multi-centre study: regional distribution and genotype-phenotype profiling of <i>SRD5A2</i> in 190 Chinese patients. <i>Journal of Medical Genetics</i> , 2019, 56, 685-692.	1.5	16
4	Phenotype and Molecular Characterizations of 30 Children From China With NR5A1 Mutations. <i>Frontiers in Pharmacology</i> , 2018, 9, 1224.	1.6	14
5	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
6	Whole Exome Sequencing Uncovered the Genetic Architecture of Growth Hormone Deficiency Patients. <i>Frontiers in Endocrinology</i> , 2021, 12, 711991.	1.5	6
7	Aromatase deficiency: A case series of 46, XX Chinese children and a systematic review of the literature. <i>Clinical Endocrinology</i> , 2020, 93, 687-695.	1.2	3
8	Exploring the efficacy of testosterone undecanoate in male children with 5 $\alpha$ -reductase deficiency. <i>Pediatric Investigation</i> , 2021, 5, 249-254.	0.6	3
9	First Clinical Study on Long-Acting Growth Hormone Therapy in Children with Turner Syndrome. <i>Hormone and Metabolic Research</i> , 2022, 54, 389-395.	0.7	2
10	SOX2 heterozygous mutations cause multiple extraocular phenotypes in boys. <i>Chinese Medical Journal</i> , 2021, Publish Ahead of Print, .	0.9	1
11	SUN-LB041 Clinical and Molecular Characteristics, Genotype-Phenotype Correlation in 113 Chinese Children with SRD5A2 Gene Mutations. <i>Journal of the Endocrine Society</i> , 2019, 3, .	0.1	0
12	Growth Curves of Chinese Children with Androgen Insensitivity Syndrome: A Multicenter Registry Study. <i>Journal of Personalized Medicine</i> , 2022, 12, 771.	1.1	0
13	DNA Hypermethylation and a Specific Methylation Spectrum on the X Chromosome in Turner Syndrome as Determined by Nanopore Sequencing. <i>Journal of Personalized Medicine</i> , 2022, 12, 872.	1.1	0