Zhiya Dong

List of Publications by Year
in descending order

15	176	1307594 7	1199594 12
papers	citations	h-index	g-index

16	16	16	280
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Clinical and molecular characterization of thirty Chinese patients with congenital lipoid adrenal hyperplasia. Journal of Steroid Biochemistry and Molecular Biology, 2021, 206, 105788.	2.5	9
2	Two lossâ€ofâ€function ANKRD11 variants in Chinese patients with short stature and a possible molecular pathway. American Journal of Medical Genetics, Part A, 2021, 185, 710-718.	1.2	7
3	A novel mutation in NF1 gene of patient with Neurofibromatosis type 1: A case report and functional study. Molecular Genetics & Genomic Medicine, 2021, 9, e1643.	1.2	1
4	The Impact of Gut Microbiome on Metabolic Disorders During Catch-Up Growth in Small-for-Gestational-Age. Frontiers in Endocrinology, 2021, 12, 630526.	3.5	7
5	Characterization of Two Loss-of-Function NF1 Variants in Chinese Patients and Potential Molecular Interpretations of Phenotypes. Frontiers in Genetics, 2021, 12, 660592.	2.3	1
6	Characterization With Gene Mutations in Han Chinese Patients With Hypospadias and Function Analysis of a Novel AR Genevariant. Frontiers in Genetics, 2021, 12, 673732.	2.3	1
7	A Multicenter Survey of Type I Diabetes Mellitus in Chinese Children. Frontiers in Endocrinology, 2021, 12, 583114.	3.5	9
8	A Novel Loss-of-Function MKRN3 Variant in a Chinese Patient With Familial Precocious Puberty: A Case Report and Functional Study. Frontiers in Genetics, 2021, 12, 663746.	2.3	4
9	Severe constipation as the first clinical manifestation in multiple endocrine neoplasia type 2B: a case report and literature review. BMC Pediatrics, 2020, 20, 318.	1.7	12
10	MKRN3 regulates the epigenetic switch of mammalian puberty via ubiquitination of MBD3. National Science Review, 2020, 7, 671-685.	9.5	48
11	Identification of gene variants in 130 Han Chinese patients with hypospadias by targeted nextâ€generation sequencing. Molecular Genetics & Genomic Medicine, 2019, 7, e827.	1.2	14
12	Pathogenic gene screening in 91 Chinese patients with short stature of unknown etiology with a targeted next-generation sequencing panel. BMC Medical Genetics, 2018, 19, 212.	2.1	18
13	17Î²-Hydroxysteroid dehydrogenase 3 deficiency: Three case reports and a systematic review. Journal of Steroid Biochemistry and Molecular Biology, 2017, 174, 141-145.	2.5	30
14	Hormone Inhibition During Mini-Puberty and Testicular Function in Male Rats. International Journal of Endocrinology and Metabolism, 2015, 13, e25465.	1.0	9
15	Mutation analysis of the SRD5A2, AR and SF-1 genes in 52 Chinese boys with hypospadias. Journal of Pediatric Endocrinology and Metabolism, 2013, 26, 887-93.	0.9	6