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	MKRN3 regulates the epigenetic switch of mammalian puberty via ubiquitination of MBD3. National Science Review, 2020, 7, 671-685.	9.5	48
2	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
3	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
4	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
5	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
6	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
7	A Multicenter Survey of Type I Diabetes Mellitus in Chinese Children. Frontiers in Endocrinology, 2021, 12, 583114.	3.5	9
8	Hormone Inhibition During Mini-Puberty and Testicular Function in Male Rats. International Journal of Endocrinology and Metabolism, 2015, 13, e25465.	1.0	9
9	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
10	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
11	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
12	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
13	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
14	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
15	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