Hui Miao

List of Publications by Year
in descending order

12	104	1478505 6	1372567 10
papers	citations	h-index	g-index

12	12	12	168
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Analysis of novel heterozygous mutations in the CYP11B2 gene causing congenital aldosterone synthase deficiency and literature review. Steroids, 2019, 150, 108448.	1.8	16
2	Clinical Characteristics of Short-Stature Patients With an NPR2 Mutation and the Therapeutic Response to rhGH. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 431-441.	3.6	15
3	Effect of 3 NR3C1 Mutations in the Pathogenesis of Pituitary ACTH Adenoma. Endocrinology, 2021, 162, .	2.8	14
4	Growth-Promoting Therapies May Be Useful In Short Stature Patients With Nonspecific Skeletal Abnormalities Caused By Acan Heterozygous Mutations: Six Chinese Cases And Literature Review. Endocrine Practice, 2020, 26, 1255-1268.	2.1	13
5	Ghrelin Promotes Proliferation and Inhibits Differentiation of 3T3-L1 and Human Primary Preadipocytes. Frontiers in Physiology, 2019, 10, 1296.	2.8	12
6	Discovery and characterization of circulating tumor cell clusters in neuroendocrine tumor patients using nanosubstrate-embedded microchips. Biosensors and Bioelectronics, 2022, 199, 113854.	10.1	10
7	A human ACTH-secreting corticotroph tumoroid model. EBioMedicine, 2021, 66, 103294.	6.1	8
8	Dwarfism in Troyer syndrome: a family with SPG20 compound heterozygous mutations and a literature review. Annals of the New York Academy of Sciences, 2020, 1462, 118-127.	3.8	7
9	A report of 2 cases of Cornelia de Lange syndrome (CdLS) and an analysis of clinical and genetic characteristics in a Chinese CdLS cohort. Molecular Genetics & Genomic Medicine, 2020, 8, e1066.	1.2	5
10	Experience of Ectopic Adrenocorticotropin Syndrome: 88 Cases With Identified Causes. Endocrine Practice, 2021, 27, 866-873.	2.1	3
11	Identification and In Vitro Functional Verification of Two Novel Mutations of GHR Gene in the Chinese Children with Laron Syndrome. Frontiers in Endocrinology, 2021, 12, 605736.	3.5	1
12	SUN-LB008 Clinical Phenotype and Genotype Analysis of Short Stature Patients with Skeletal Dysplasia. Journal of the Endocrine Society, 2019, 3, .	0.2	0