Chao-Chun Zou

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/4629985/publications.pdf

Version: 2024-02-01

15 papers	101 citations	1478505 6 h-index	1474206 9 g-index
20 all docs	20 docs citations	20 times ranked	89 citing authors

#	Article	IF	Citations
1	Regulatory roles and mechanisms of alternative RNA splicing in adipogenesis and human metabolic health. Cell and Bioscience, 2021, 11, 66.	4.8	26
2	Application of next generation sequencing in the screening of monogenic diseases in China, 2021: a consensus among Chinese newborn screening experts. World Journal of Pediatrics, 2022, 18, 235-242.	1.8	16
3	Glucose metabolism disorder in obese children assessed by continuous glucose monitoring system. World Journal of Pediatrics, 2008, 4, 26-30.	1.8	13
4	Clinical features and genetic analysis of childhood sitosterolemia. Medicine (United States), 2019, 98, e15013.	1.0	13
5	Quality of life in caregivers of young children with Praderâ€“Willi syndrome. World Journal of Pediatrics, 2019, 15, 506-510.	1.8	9
6	Recommendations for the diagnosis and management of childhood Prader-Willi syndrome in China. Orphanet Journal of Rare Diseases, 2022, 17, .	2.7	8
7	Prader-Willi Syndrome: Molecular Mechanism and Epigenetic Therapy. Current Gene Therapy, 2020, 20, 36-43.	2.0	7
8	Do patients with Praderâ€“Willi syndrome have favorable glucose metabolism?. Orphanet Journal of Rare Diseases, 2022, 17, 187.	2.7	4
9	Clinical Features in Patients with Microdeletion at 6q14.1-q15. Indian Journal of Pediatrics, 2017, 84, 883-886.	0.8	1
10	Prader-Willi Syndrome Coincident with DiGeorge Syndrome. Indian Journal of Pediatrics, 2020, 87, 471-472.	0.8	1
11	6q25.1-q25.3 Microdeletion in a Chinese Girl. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2021, 13, 109-113.	0.9	1
12	13q22.1-q32.1 Microdeletion Syndrome. Indian Journal of Pediatrics, 2019, 86, 303-305.	0.8	0
13	Clinical Features of Patients With 7p22.1 Microdeletion. Indian Pediatrics, 2020, 57, 580-581.	0.4	0
14	Clinical Features of Patients With 7p22.1 Microdeletion. Indian Pediatrics, 2020, 57, 580-581.	0.4	0
15	Microdeletion of 4p16.2 in Children: A Case Report and Literature Review. Case Reports in Genetics, 2022, 2022, 1-4.	0.2	0