

Zhou Yang

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3826973/publications.pdf>

Version: 2024-02-01

11
papers

92
citations

1936888

4
h-index

1372195

10
g-index

11
all docs

11
docs citations

11
times ranked

198
citing authors

#	ARTICLE	IF	CITATIONS
1	Characterization of community-associated <i>Staphylococcus aureus</i> from skin and soft-tissue infections: a multicenter study in China. <i>Emerging Microbes and Infections</i> , 2016, 5, 1-11.	3.0	36
2	Peripheral refraction in 7- and 14-year-old children in central China: the Anyang Childhood Eye Study. <i>British Journal of Ophthalmology</i> , 2015, 99, 674-679.	2.1	26
3	Heterozygous somatic activating AKT1 mutation in a case of Proteus syndrome with mental retardation. <i>Journal of Dermatology</i> , 2014, 41, 188-189.	0.6	6
4	Large Deletions in the NSDHL Gene in Two Patients with CHILD Syndrome. <i>Acta Dermato-Venereologica</i> , 2015, 95, 1007-1008.	0.6	6
5	Toxic epidermal necrolysis after dactinomycin and vincristine combination chemotherapy for nephroblastoma. <i>Journal of Zhejiang University: Science B</i> , 2017, 18, 649-652.	1.3	5
6	SERPINB7 novel mutation in Chinese patients with Nagashima-type palmoplantar keratosis and cases associated with atopic dermatitis. <i>International Journal of Dermatology</i> , 2020, 59, e320-e322.	0.5	4
7	Application of topical gentamicin—a new era in the treatment of genodermatosis. <i>World Journal of Pediatrics</i> , 2021, 17, 568-575.	0.8	3
8	A novel frameshift truncation mutation in the V2 tail domain of KRT1 causes mild ichthyosis hystrix of Curth-Macklin. <i>Clinical and Experimental Dermatology</i> , 2020, 45, 719-721.	0.6	2
9	Abnormal resting-state brain activity and connectivity of brain-bladder control network in overactive bladder syndrome. <i>Acta Radiologica</i> , 2022, 63, 1695-1702.	0.5	2
10	Novel <i>MBTPS2</i> mutation causes a mild phenotype of ichthyosis follicularis with atrichia and photophobia syndrome in a Chinese pedigree. <i>Journal of Dermatology</i> , 2019, 46, e126-e128.	0.6	1
11	Novel ABCA12 compound heterozygous mutations identified in a patient with congenital ichthyosiform erythroderma and aortopulmonary window. <i>European Journal of Dermatology</i> , 2019, 29, 83-85.	0.3	1