Hong Yu

List of Publications by Year
in descending order

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14 papers	120 citations	1937685 4 h-index	1281871 11 g-index
15 all docs	15 docs citations	15 times ranked	176 citing authors

#	Article	IF	Citations
1	Woolly hair nevus caused by somatic mutation and Costello syndrome caused by germline mutation in HRAS: Consider parental mosaicism in prenatal counseling. Journal of Dermatology, 2022, 49, 161-164.	1.2	3
2	Facial Balamuthia mandrillaris infection with neurological involvement in an immunocompetent child. Lancet Infectious Diseases, The, 2022, 22, e93-e100.	9.1	4
3	Majocchi's granuloma on the scalp of a 55â€yearâ€old female successfully treated with terbinafine. Dermatologic Therapy, 2022, , e15304.	1.7	0
4	Damaged Keratin Filament Network Caused by KRT5 Mutations in Localized Recessive Epidermolysis Bullosa Simplex. Frontiers in Genetics, 2021, 12, 736610.	2.3	3
5	Cutaneous metastases from triple primary extramammary Pagetâ€™s disease. JDDG - Journal of the German Society of Dermatology, 2020, 18, 1169-1172.	0.8	3
6	Excellent response to oral clarithromycin in a patient with severe childhood granulomatous periorificial dermatitis with neck involvement. Journal of Dermatology, 2020, 47, e222-e224.	1.2	4
7	Painful Indurated Plaques on the Vulva. JAMA Dermatology, 2019, 155, 1073.	4.1	0
8	Severe dermatitis, multiple allergies and metabolic wasting (<scp>SAM</scp>) syndrome caused by de novo mutation in the DSP gene misdiagnosed as generalized pustular psoriasis and treatment of acitretin with gabapentin. Journal of Dermatology, 2019, 46, 622-625.	1.2	14
9	Ankyloblepharonâ€“ectodermal dysplasiaâ€“clefting syndrome misdiagnosed as epidermolysis bullosa and congenital ichthyosiform erythroderma: Case report and review of published work. Journal of Dermatology, 2019, 46, 422-425.	1.2	11
10	Generalized eruptive keratoacanthoma with vitiligo followed by the development of prurigo nodularis: A case report and published work review. Journal of Dermatology, 2018, 45, 211-215.	1.2	1
11	Typical Chinese pedigree of autosomal dominant genetic disease: Neurofibromatosis type 1 with a novel frameâ€shift mutation. Journal of Dermatology, 2017, 44, e188-e189.	1.2	0
12	H19 lncRNA regulates keratinocyte differentiation by targeting miR-130b-3p. Cell Death and Disease, 2017, 8, e3174-e3174.	6.3	52
13	Genome Sequence of a Novel Recombinant Coxsackievirus A6 Strain from Shanghai, China, 2013. Genome Announcements, 2015, 3, .	0.8	3
14	A child with coexistent juvenile xanthogranuloma and Langerhans cell histiocytosis. Journal of the American Academy of Dermatology, 2010, 62, 329-332.	1.2	21