

Zhirong Yao

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

79
papers

729
citations

14
h-index

24
g-index

85
ext. papers

977
ext. citations

3.8
avg, IF

3.69
L-index

#	Paper	IF	Citations
79	Majocchi's granuloma on the scalp of a 55-year-old female successfully treated with terbinafine.. <i>Dermatologic Therapy</i> , 2022 , e15304	2.2	
78	Segmental vitiligo following acitretin treatment for infantile generalized pustular psoriasis resulting in repigmentation under secukinumab therapy.. <i>Dermatologic Therapy</i> , 2022 , e15305	2.2	
77	The clinical efficacy and safety of anti-IgE therapy in recessive dystrophic epidermolysis bullosa. <i>Clinical Genetics</i> , 2022 , 101, 110-115	4	1
76	Complex genetic models in dystrophic epidermolysis bullosa families with marked intra-familial phenotypic heterogeneity.. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2022	4.6	0
75	Detection and characterization of low-level mosaicism among clinically unaffected parents of "sporadic" epidermolysis bullosa simplex cases.. <i>British Journal of Dermatology</i> , 2022 ,	4	0
74	S1P defects cause a new entity of cataract, alopecia, oral mucosal disorder, and psoriasis-like syndrome.. <i>EMBO Molecular Medicine</i> , 2022 , e14904	12	1
73	Damaged Keratin Filament Network Caused by Mutations in Localized Recessive Epidermolysis Bullosa Simplex.. <i>Frontiers in Genetics</i> , 2021 , 12, 736610	4.5	1
72	Woolly hair nevus caused by somatic mutation and Costello syndrome caused by germline mutation in HRAS: Consider parental mosaicism in prenatal counseling. <i>Journal of Dermatology</i> , 2021 , 49, 161	1.6	2
71	Deep-intronic and frameshift DSG1 variants associated with atypical severe dermatitis, multiple allergies and metabolic wasting (SAM) syndrome in a Chinese family. <i>European Journal of Dermatology</i> , 2021 , 31, 239-244	0.8	0
70	Additional causal SNRPE mutations in hereditary hypotrichosis simplex. <i>British Journal of Dermatology</i> , 2021 , 185, 439-441	4	1
69	Factors associated with persistence of early-onset atopic dermatitis up to the age of 12 years: a prospective cohort study in China. <i>European Journal of Dermatology</i> , 2021 , 31, 403-408	0.8	1
68	Identification of a T 2-high psoriasis cluster based on skin biomarker analysis in a Chinese psoriasis population. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2021 , 35, 150-158	4.6	3
67	Simvastatin ointment in the treatment of seven childhood diffuse plane xanthomas. <i>Journal of Dermatology</i> , 2021 , 48, 223-227	1.6	1
66	Randomized clinical trial of combined therapy with oral lipoic acid and NB-UVB for nonsegmental stable vitiligo. <i>Dermatologic Therapy</i> , 2021 , 34, e14610	2.2	1
65	Netherton syndrome caused by compound heterozygous mutation, c.80A>G mutation in SPINK5 and large-sized genomic deletion mutation, and successful treatment of intravenous immunoglobulin. <i>Molecular Genetics & Genomic Medicine</i> , 2021 , 9, e1600	2.3	2
64	A Case of Familial Cold Autoinflammatory Syndrome with Mutation. <i>Annals of Dermatology</i> , 2021 , 33, 198-200	0.4	
63	Roles of the H19/microRNA-675 axis in the proliferation and epithelial-mesenchymal transition of human cutaneous squamous cell carcinoma cells. <i>Oncology Reports</i> , 2021 , 45,	3.5	5

62	Update on the Pathogenesis and Therapy of Atopic Dermatitis. <i>Clinical Reviews in Allergy and Immunology</i> , 2021 , 1	12.3	10
61	Efficacy and Safety of Abrocitinib in Combination With Topical Therapy in Adolescents With Moderate-to-Severe Atopic Dermatitis: The JADE TEEN Randomized Clinical Trial. <i>JAMA Dermatology</i> , 2021 , 157, 1165-1173	5.1	11
60	Comparison of the efficacy and safety between a low-fluence 1064-nm Q-switched neodymium-doped yttrium aluminum garnet laser and a conventional Q-switched 532-nm laser for the treatment of cafe-au-lait macules in 40 Chinese children: a prospective, randomized, parallel-controlled, evaluator-blinded trial. <i>Lasers in Medical Science</i> , 2021 , 1	3.1	1
59	Early Life Domestic Pet Ownership, and the Risk of Pet Sensitization and Atopic Dermatitis in Preschool Children: A Prospective Birth Cohort in Shanghai. <i>Frontiers in Pediatrics</i> , 2020 , 8, 192	3.4	4
58	Next-generation sequencing through multigene panel testing for the diagnosis of hereditary epidermolysis bullosa in Chinese population. <i>Clinical Genetics</i> , 2020 , 98, 179-184	4	4
57	Smad2/4 Signaling Pathway Is Critical for Epidermal Langerhans Cell Repopulation Under Inflammatory Condition but Not Required for Their Homeostasis at Steady State. <i>Frontiers in Immunology</i> , 2020 , 11, 912	8.4	1
56	An Indurated Plaque on the Eyelid. <i>JAMA Dermatology</i> , 2020 , 156, 811-812	5.1	
55	Excellent response to oral clarithromycin in a patient with severe childhood granulomatous periorificial dermatitis with neck involvement. <i>Journal of Dermatology</i> , 2020 , 47, e222-e224	1.6	1
54	Case of acne-like lesion of cutaneous candidiasis in a healthy man. <i>International Journal of Dermatology</i> , 2020 , 59, e148-e150	1.7	
53	Cronkhite-Canada syndrome in an adult with titanium orthopaedic implants. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2020 , 34, e315-e317	4.6	1
52	Gentamicin induces COL17A1 nonsense mutation readthrough in junctional epidermolysis bullosa. <i>Journal of Dermatology</i> , 2020 , 47, e82-e83	1.6	9
51	A novel MBTPS2 start codon mutation causes a mild ichthyosis follicularis with atrichia and photophobia phenotype. <i>Clinical and Experimental Dermatology</i> , 2020 , 45, 505-507	1.8	0
50	Concurrence of generalized perforating and subcutaneous granuloma annulare in a 4-year-old boy with latent tuberculosis infection successfully treated with low-dose hydroxychloroquine. <i>Journal of Dermatology</i> , 2020 , 47, e71-e72	1.6	2
49	Cutaneous metastases from triple primary extramammary Paget disease. <i>JDDG - Journal of the German Society of Dermatology</i> , 2020 , 18, 1169-1172	1.2	1
48	The first case of a mosaic superficial epidermolytic ichthyosis diagnosed by Ultra-Deep Sequence. <i>Molecular Genetics & Genomic Medicine</i> , 2020 , 8, e1457	2.3	4
47	Development and validation of new diagnostic criteria for atopic dermatitis in children of China. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2020 , 34, 542-548	4.6	5
46	An excellent response to topical therapy of four congenital hemidysplasia with ichthyosiform erythroderma and limb defects syndrome patients with an increased concentration of simvastatin ointment. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2020 , 34, e8-e11	4.6	2
45	Severe dermatitis, multiple allergies and metabolic wasting (SAM) syndrome caused by de novo mutation in the DSP gene misdiagnosed as generalized pustular psoriasis and treatment of acitretin with gabapentin. <i>Journal of Dermatology</i> , 2019 , 46, 622-625	1.6	7

44	Mutation in FAM111B Causes Hereditary Fibrosing Poikiloderma with Tendon Contracture, Myopathy, and Pulmonary Fibrosis. <i>Acta Dermato-Venereologica</i> , 2019 , 99, 695-696	2.2	2
43	Phenotypic analysis of atopic dermatitis in children aged 1-12 months: elaboration of novel diagnostic criteria for infants in China and estimation of prevalence. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2019 , 33, 1569-1576	4.6	12
42	Family of hereditary fibrosing poikiloderma with tendon contractures, myopathy and pulmonary fibrosis caused by a novel FAM111B mutation. <i>Journal of Dermatology</i> , 2019 , 46, 1014-1018	1.6	1
41	Coexistence of hypopigmented mycosis fungoides and erythema dyschromicum perstans in a 3-year-old Chinese girl. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2019 , 33, e492-e494	4.6	494
40	Painful Indurated Plaques on the Vulva. <i>JAMA Dermatology</i> , 2019 , 155, 1073-1074	5.1	
39	Coinheritance of generalized pustular psoriasis and familial Behçet-like autoinflammatory syndrome with variants in IL36RN and TNFAIP3 in the heterozygous state. <i>Journal of Dermatology</i> , 2019 , 46, 907-910	1.6	5
38	Three novel mutations in GPNMB in two pedigrees with amyloidosis cutis dyschromica. <i>British Journal of Dermatology</i> , 2019 , 181, 1327-1329	4	8
37	Ankyloblepharon-ectodermal dysplasia-clefting syndrome misdiagnosed as epidermolysis bullosa and congenital ichthyosiform erythroderma: Case report and review of published work. <i>Journal of Dermatology</i> , 2019 , 46, 422-425	1.6	8
36	CHILD syndrome mimicking verrucous nevus in a Chinese patient responded well to the topical therapy of compound of simvastatin and cholesterol. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2018 , 32, 1209-1213	4.6	6
35	Generalized eruptive keratoacanthoma with vitiligo followed by the development of prurigo nodularis: A case report and published work review. <i>Journal of Dermatology</i> , 2018 , 45, 211-215	1.6	1
34	A Missense Mutation within the Helix Termination Motif of KRT25 Causes Autosomal Dominant Woolly Hair/Hypotrichosis. <i>Journal of Investigative Dermatology</i> , 2018 , 138, 230-233	4.3	5
33	LUMBAR syndrome: A case manifesting as cutaneous infantile hemangiomas of the lower extremity, perineum and gluteal region, and a review of published work. <i>Journal of Dermatology</i> , 2017 , 44, 808-812	1.6	6
32	Mutations in IL36RN are associated with geographic tongue. <i>Human Genetics</i> , 2017 , 136, 241-252	6.3	24
31	Skin involvement as the first symptom of rapidly progressive ALK-positive systemic anaplastic large cell lymphoma. <i>Clinical and Experimental Dermatology</i> , 2017 , 42, 539-542	1.8	2
30	Expansion of the genotypic and phenotypic spectrum of xeroderma pigmentosum in Chinese population. <i>Photodermatology Photoimmunology and Photomedicine</i> , 2017 , 33, 58-63	2.4	4
29	Clinical profiles of pediatric patients with GPP alone and with different IL36RN genotypes. <i>Journal of Dermatological Science</i> , 2017 , 85, 235-240	4.3	12
28	A novel mutation in TRPV3 gene causes atypical familial Olmsted syndrome. <i>Scientific Reports</i> , 2016 , 6, 21815	4.9	24
27	Identification of a PTPN11 hot spot mutation in a child with atypical LEOPARD syndrome. <i>Molecular Medicine Reports</i> , 2016 , 14, 2639-43	2.9	5

26	Report of Chinese family with severe dermatitis, multiple allergies and metabolic wasting syndrome caused by novel homozygous desmoglein-1 gene mutation. <i>Journal of Dermatology</i> , 2016 , 43, 1201-1204	1.6	14
25	First Mal de Meleda report in Chinese Mainland: two families with a recurrent homozygous missense mutation in SLURP-1. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2016 , 30, 871-3	4.6	8
24	Effects of lidocaine on regulatory T cells in atopic dermatitis. <i>Journal of Allergy and Clinical Immunology</i> , 2016 , 137, 613-617.e5	11.5	10
23	Genotype analysis of varicella-zoster virus isolates from suburban Shanghai Municipal Province, China. <i>Journal of Medical Microbiology</i> , 2016 , 65, 123-128	3.2	3
22	Genome-wide linkage and exome sequencing analyses identify an initiation codon mutation of KRT5 in a unique Chinese family with generalized Dowling-Degos disease. <i>British Journal of Dermatology</i> , 2016 , 174, 663-6	4	6
21	Lentiginous phenotypes caused by diverse pathogenic genes (SASH1 and PTPN11): clinical and molecular discrimination. <i>Clinical Genetics</i> , 2016 , 90, 372-7	4	13
20	A rare RECQL4 indel mutation in a Chinese patient with Rothmund-Thomson syndrome. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2016 , 30, e159-e161	4.6	2
19	Gene diagnosis and prenatal genetic diagnosis of a case of dystrophic epidermolysis bullosa family caused by gonadosomatic mosaicism for the COL7A1 mutation p.Gly2043Arg in the pregnant mother. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2016 , 30, 1627-9	4.6	2
18	Report of a child with sporadic familial progressive hyper- and hypopigmentation caused by a novel KITLG mutation. <i>British Journal of Dermatology</i> , 2016 , 175, 1369-1371	4	9
17	Prevalence of Atopic Dermatitis in Chinese Children aged 1-7 ys. <i>Scientific Reports</i> , 2016 , 6, 29751	4.9	57
16	Genome-wide linkage analysis and whole-genome sequencing identify a recurrent SMARCAD1 variant in a unique Chinese family with Basan syndrome. <i>European Journal of Human Genetics</i> , 2016 , 24, 1367-70	5.3	12
15	Novel MBTPS2 missense mutation causes a keratosis follicularis spinulosa decalvans phenotype: mutation update and review of the literature. <i>Clinical and Experimental Dermatology</i> , 2016 , 41, 757-60	1.8	10
14	Mutations in the mevalonate pathway genes in Chinese patients with porokeratosis. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2016 , 30, 1512-7	4.6	14
13	Molecular Characterization of NF1 and Neurofibromatosis Type 1 Genotype-Phenotype Correlations in a Chinese Population. <i>Scientific Reports</i> , 2015 , 5, 11291	4.9	37
12	Genome sequence of a novel recombinant coxsackievirus a6 strain from shanghai, china, 2013. <i>Genome Announcements</i> , 2015 , 3,		3
11	Development of two molecular approaches for differentiation of clinically relevant yeast species closely related to <i>Candida guilliermondii</i> and <i>Candida famata</i> . <i>Journal of Clinical Microbiology</i> , 2014 , 52, 3190-5	9.7	7
10	Prevalent and rare mutations in IL-36RN gene in Chinese patients with generalized pustular psoriasis and psoriasis vulgaris. <i>Journal of Investigative Dermatology</i> , 2013 , 133, 2637-2639	4.3	47
9	Simultaneous identification of molecular and mating types within the <i>Cryptococcus</i> species complex by PCR-RFLP analysis. <i>Journal of Medical Microbiology</i> , 2008 , 57, 1481-1490	3.2	26

8	Fungal respiratory disease. <i>Current Opinion in Pulmonary Medicine</i> , 2006 , 12, 222-7	3	28
7	Management of cryptococcosis in non-HIV-related patients. <i>Medical Mycology</i> , 2005 , 43, 245-51	3.9	27
6	The relationship of glycosylation and isoelectric point with tumor accumulation of avidin. <i>Journal of Nuclear Medicine</i> , 1999 , 40, 479-83	8.9	33
5	Imaging of intraperitoneal tumors with technetium-99m GSA. <i>Annals of Nuclear Medicine</i> , 1998 , 12, 115-8.5	8.5	17
4	Increased streptavidin uptake in tumors pretargeted with biotinylated antibody using a conjugate of streptavidin-fab fragment. <i>Nuclear Medicine and Biology</i> , 1998 , 25, 557-60	2.1	3
3	Avidin targeting of intraperitoneal tumor xenografts. <i>Journal of the National Cancer Institute</i> , 1998 , 90, 25-9	9.7	74
2	Radioimmunoimaging of colon cancer xenografts with anti-Tn monoclonal antibody. <i>Nuclear Medicine and Biology</i> , 1995 , 22, 199-203	2.1	10
1	Improved targeting of radiolabeled streptavidin in tumors pretargeted with biotinylated monoclonal antibodies through an avidin chase. <i>Journal of Nuclear Medicine</i> , 1995 , 36, 837-41	8.9	44