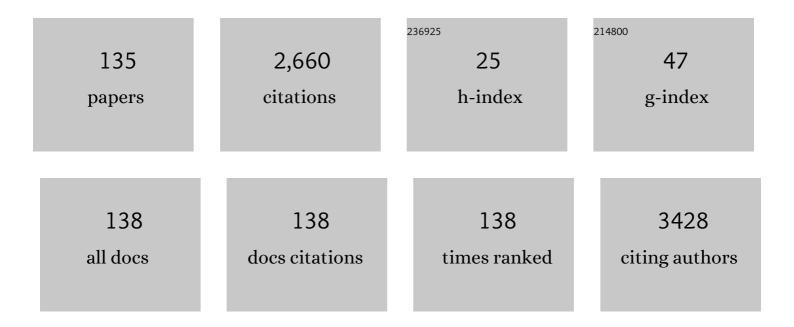
Chao-Kai Hsu

List of Publications by Year in descending order

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Сило-Клі Нец

#	Article	IF	CITATIONS
1	Genetic Variants Associated With Phenytoin-Related Severe Cutaneous Adverse Reactions. JAMA - Journal of the American Medical Association, 2014, 312, 525.	7.4	256
2	Erythema Multiforme, Stevens-Johnson Syndrome, and Toxic Epidermal Necrolysis. Cornea, 2007, 26, 123-129.	1.7	159
3	Demodicosis: A clinicopathological study. Journal of the American Academy of Dermatology, 2009, 60, 453-462.	1.2	152
4	Risk and association of <i>HLA</i> with oxcarbazepine-induced cutaneous adverse reactions in Asians. Neurology, 2017, 88, 78-86.	1.1	117
5	Lichen planus and lichenoid dermatoses. Journal of the American Academy of Dermatology, 2018, 79, 789-804.	1.2	116
6	The tension biology of wound healing. Experimental Dermatology, 2019, 28, 464-471.	2.9	116
7	Paper-Based ELISA for the Detection of Autoimmune Antibodies in Body Fluid—The Case of Bullous Pemphigoid. Analytical Chemistry, 2014, 86, 4605-4610.	6.5	90
8	Mechanical forces in skin disorders. Journal of Dermatological Science, 2018, 90, 232-240.	1.9	78
9	Caveolin-1 Controls Hyperresponsiveness to Mechanical Stimuli and Fibrogenesis-Associated RUNX2 ActivationÂin Keloid Fibroblasts. Journal of Investigative Dermatology, 2018, 138, 208-218.	0.7	74
10	Successful Treatment of Recalcitrant Pemphigus Vulgaris and Pemphigus Vegetans With Etanercept and Carbon Dioxide Laser. Archives of Dermatology, 2005, 141, 680.	1.4	69
11	Evanescent and Persistent Pruritic Eruptions of Adult-Onset Still Disease: A Clinical and Pathologic Study of 36 Patients. Seminars in Arthritis and Rheumatism, 2012, 42, 317-326.	3.4	64
12	Mechanobiological dysregulation of the epidermis and dermis in skin disorders and in degeneration. Journal of Cellular and Molecular Medicine, 2013, 17, 817-822.	3.6	61
13	The Medication Risk of Stevens–Johnson Syndrome and Toxic Epidermal Necrolysis in Asians: The Major Drug Causality and Comparison With the US FDA Label. Clinical Pharmacology and Therapeutics, 2019, 105, 112-120.	4.7	54
14	Analysis of Taiwanese ichthyosis vulgaris families further demonstrates differences in <i>FLG</i> mutations between European and Asian populations. British Journal of Dermatology, 2009, 161, 448-451.	1.5	49
15	Higher body mass index is associated with greater severity of alopecia in men with male-pattern androgenetic alopecia in Taiwan: A cross-sectional study. Journal of the American Academy of Dermatology, 2014, 70, 297-302.e1.	1.2	49
16	Treatment of Hereditary Epidermolysis Bullosa: Updates and Future Prospects. American Journal of Clinical Dermatology, 2014, 15, 1-6.	6.7	46
17	Whole genome sequencing identifies genetic variants associated with co-trimoxazole hypersensitivity in Asians. Journal of Allergy and Clinical Immunology, 2021, 147, 1402-1412.	2.9	46
18	Time Series Integrative Analysis of RNA SequencingÂand MicroRNA Expression Data RevealsÂKey Biologic Wound Healing PathwaysÂinÂKeloid-Prone Individuals. Journal of Investigative Dermatology, 2018, 138, 2690-2693.	0.7	41

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#	Article	IF	CITATIONS
19	Use of hydrochlorothiazide and risk of skin cancer: a nationwide Taiwanese case–control study. British Journal of Cancer, 2019, 121, 973-978.	6.4	40
20	Beneficial effect of ustekinumab in familial pityriasis rubra pilaris with a new missense mutation in <i>CARD14</i> . British Journal of Dermatology, 2018, 178, 969-972.	1.5	38
21	Mutations in KLHL24 Add to the Molecular Heterogeneity of Epidermolysis BullosaÂSimplex. Journal of Investigative Dermatology, 2017, 137, 1378-1380.	0.7	37
22	Noninvasive evaluation of collagen and hemoglobin contents and scattering property of <i>in vivo</i> keloid scars and normal skin using diffuse reflectance spectroscopy: pilot study. Journal of Biomedical Optics, 2012, 17, 0770051.	2.6	36
23	Non-invasive evaluation of therapeutic response in keloid scar using diffuse reflectance spectroscopy. Biomedical Optics Express, 2015, 6, 390.	2.9	36
24	Mechanical coupling of cytoskeletal elasticity and force generation is crucial for understanding the migrating nature of keloid fibroblasts. Experimental Dermatology, 2015, 24, 579-584.	2.9	32
25	Large Intragenic Deletion in DSTYK Underlies Autosomal-Recessive Complicated Spastic Paraparesis, SPG23. American Journal of Human Genetics, 2017, 100, 364-370.	6.2	32
26	Linear lupus panniculitis of the scalp presenting as alopecia along Blaschko's lines: A distinct variant of lupus panniculitis in East Asians?. Journal of Dermatology, 2012, 39, 385-388.	1.2	28
27	Dissecting folliculitis (dissecting cellulitis) of the scalp: a 66â€patient case series and proposal of classification. JDDG - Journal of the German Society of Dermatology, 2018, 16, 1219-1226.	0.8	26
28	Investigational Treatments for Epidermolysis Bullosa. American Journal of Clinical Dermatology, 2021, 22, 801-817.	6.7	26
29	Clinicopathological study of Fox–Fordyce disease. Journal of Dermatology, 2009, 36, 485-490.	1.2	24
30	Progressive hyperpigmentation in a Taiwanese child due to an inborn error of vitamin B12 metabolism (cblJ). British Journal of Dermatology, 2015, 172, 1111-1115.	1.5	24
31	Congenital Anonychia and Uncombable Hair Syndrome: Coinheritance of Homozygous Mutations in RSPO4 and PADI3. Journal of Investigative Dermatology, 2017, 137, 1176-1179.	0.7	23
32	Lichen planus and lichenoid dermatoses. Journal of the American Academy of Dermatology, 2018, 79, 807-818.	1.2	23
33	Exacerbation of psoriatic skin lesions in a patient with Alzheimer disease receiving gamma-secretase inhibitor. Journal of the American Academy of Dermatology, 2013, 68, e46-e48.	1.2	22
34	Syndromic inherited poikiloderma due to a <i>de novo</i> mutation in <i> <scp>FAM</scp> 111B </i> . British Journal of Dermatology, 2017, 176, 534-536.	1.5	22
35	Thrombomodulin Promotes Diabetic Wound Healing by Regulating Toll-Like Receptor 4 Expression. Journal of Investigative Dermatology, 2015, 135, 1668-1675.	0.7	19
36	Tumor Endothelial Marker 1 (TEM1/Endosialin/CD248) Enhances Wound Healing byÂInteracting with Platelet-Derived Growth Factor Receptors. Journal of Investigative Dermatology, 2019, 139, 2204-2214.e7.	0.7	18

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#	Article	IF	CITATIONS
37	Novel homozygous missense mutation in <i>NT5C2</i> underlying hereditary spastic paraplegia SPG45. American Journal of Medical Genetics, Part A, 2017, 173, 3109-3113.	1.2	17
38	Primary scarring alopecia: A retrospective study of 89 patients in Taiwan. Journal of Dermatology, 2018, 45, 450-455.	1.2	16
39	Early Intervention with High-Dose Steroid Pulse Therapy Prolongs Disease-Free Interval of Severe Alopecia Areata: A Retrospective Study. Annals of Dermatology, 2013, 25, 471.	0.9	15
40	Portable handheld diffuse reflectance spectroscopy system for clinical evaluation of skin: a pilot study in psoriasis patients. Biomedical Optics Express, 2016, 7, 616.	2.9	15
41	Homozygous acceptor splice site mutation in DSG1 disrupts plakoglobin localization and results in keratoderma and skin fragility. Journal of Dermatological Science, 2018, 89, 198-201.	1.9	14
42	Prognostic role of tumoral PDL1 expression and peritumoral FoxP3+ lymphocytes in vulvar melanomas. Human Pathology, 2018, 73, 176-183.	2.0	14
43	Adjuvant Radiotherapy After Keloid Excision. Annals of Plastic Surgery, 2019, 82, S39-S44.	0.9	14
44	Rapid detection of dermatophytes and Candida albicansin onychomycosis specimens by an oligonucleotide array. BMC Infectious Diseases, 2014, 14, 581.	2.9	13
45	Incontinentia pigmenti in a father and daughter. British Journal of Dermatology, 2016, 175, 1059-1060.	1.5	12
46	Topical betaxolol for treating relapsing paronychia with pyogenic granuloma-like lesions induced by epidermal growth factor receptor inhibitors. Journal of the American Academy of Dermatology, 2018, 78, e143-e144.	1.2	12
47	Semidominant CPNMB Mutations in Amyloidosis Cutis Dyschromica. Journal of Investigative Dermatology, 2019, 139, 2550-2554.e9.	0.7	12
48	A new superficial needleâ€scraping method for assessing <i>Demodex</i> density in papulopustular rosacea. Journal of Cosmetic Dermatology, 2020, 19, 896-900.	1.6	12
49	Coagulation Factor XIII-A Subunit Missense Mutation in the Pathobiology of Autosomal Dominant Multiple Dermatofibromas. Journal of Investigative Dermatology, 2020, 140, 624-635.e7.	0.7	12
50	Erythema Multiforme-Like Secondary Syphilis in a HIV-positive Bisexual Man. Acta Dermato-Venereologica, 2010, 90, 647-648.	1.3	11
51	A Novel Pathogenic HSPG2 Mutation in Schwartz–Jampel Syndrome. Frontiers in Neurology, 2021, 12, 632336.	2.4	11
52	Lip verrucous carcinoma in a pregnant woman successfully treated with carbon dioxide laser surgery. British Journal of Dermatology, 2007, 157, 813-815.	1.5	10
53	PLACK syndrome resulting from a new homozygous insertion mutation in CAST. Journal of Dermatological Science, 2017, 88, 256-258.	1.9	10
54	Recalcitrant extragenital giant condyloma acuminatum: A need for combination therapy. Dermatologic Therapy, 2019, 32, e12867.	1.7	10

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#	Article	IF	CITATIONS
55	Real-world efficacy of biological agents in moderate-to-severe plaque psoriasis: An analysis of 75 patients in Taiwan. PLoS ONE, 2020, 15, e0244620.	2.5	10
56	Fusariosis occurring in an ulcerated cutaneous CD8+ T cell lymphoma tumor. European Journal of Dermatology, 2006, 16, 297-301.	0.6	10
57	Current topics in Epidermolysis bullosa: Pathophysiology and therapeutic challenges. Journal of Dermatological Science, 2021, 104, 164-176.	1.9	10
58	Good cosmesis of a large rhinophyma after carbon dioxide laser treatment. Journal of Dermatology, 2006, 33, 227-229.	1.2	8
59	Large, papillomatous and pedunculated nevus sebaceus. Journal of Dermatology, 2011, 38, 200-202.	1.2	8
60	Ichthyosis Prematurity Syndrome. JAMA Dermatology, 2016, 152, 1055.	4.1	8
61	Predictive phenotyping of inherited ichthyosis by nextâ€generation <scp>DNA</scp> sequencing. British Journal of Dermatology, 2017, 176, 249-251.	1.5	8
62	"Spade sign―and inflammation/fibrosis limited to the upper and midâ€dermis as the pathognomonic features of acne keloidalis. Journal of Dermatology, 2020, 47, 41-46.	1.2	8
63	Thumbnailâ€squeezing method: an effective method for assessing Demodex density in rosacea. Journal of the European Academy of Dermatology and Venereology, 2020, 34, e343-e345.	2.4	8
64	Atypical fibroxanthoma-like amelanotic malignant melanoma: A case report and literature review. Dermatologica Sinica, 2013, 31, 140-144.	0.5	7
65	Further evidence for genotype-phenotype disparity in Griscelli syndrome. British Journal of Dermatology, 2017, 176, 1086-1089.	1.5	7
66	Two Cases of Interleukin-7–Deficient Generalized Verrucosis. Clinical Infectious Diseases, 2020, 71, 1561-1563.	5.8	7
67	Multidisciplinary care of epidermolysis bullosa during the COVID-19 pandemic—Consensus: Recommendations by an international panel of experts. Journal of the American Academy of Dermatology, 2020, 83, 1222-1224.	1.2	7
68	A germline mutation in the plateletâ€derived growth factor receptor beta gene may be implicated in hereditary progressive mucinous histiocytosis. British Journal of Dermatology, 2021, 184, 967-970.	1.5	7
69	Rosacea with persistent facial erythema and high Demodex density effectively treated with topical ivermectin alone or combined with oral carvedilol. Dermatologic Therapy, 2021, 34, e14899.	1.7	7
70	Topical gentamicin ointment induces <i>LAMB3</i> nonsense mutation readthrough and improves corneal erosions in a patient with junctional epidermolysis bullosa. Clinical and Experimental Ophthalmology, 2021, 49, 309-312.	2.6	7
71	ASC-J9 Blocks Cell Proliferation and Extracellular Matrix Production of Keloid Fibroblasts through Inhibiting STAT3 Signaling. International Journal of Molecular Sciences, 2022, 23, 5549.	4.1	7
72	Transient zinc deficiency syndrome in a breast-fed infant due to decreased zinc in breast milk (type II) Tj ETQq0 0	0 rgBT /0 0.5	Overlock 10 Tf 6

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#	Article	IF	CITATIONS
73	Broadband absorption and reduced scattering spectra of in-vivo skin can be noninvasively determined using δ-P_1 approximation based spectral analysis. Biomedical Optics Express, 2015, 6, 443.	2.9	6
74	Toward reliable retrieval of functional information of papillary dermis using spatially resolved diffuse reflectance spectroscopy. Biomedical Optics Express, 2016, 7, 542.	2.9	6
75	Spatial distribution of filament elasticity determines the migratory behaviors of a cell. Cell Adhesion and Migration, 2016, 10, 368-377.	2.7	6
76	Cutaneous lymphomas in Taiwan: A review of 118 cases from a medical center in southern Taiwan. Dermatologica Sinica, 2018, 36, 16-24.	0.5	6
77	Lateâ€onset comedonal Darier's disease caused by a recurrent <i>ATP2A2</i> mutation. Journal of Dermatology, 2019, 46, e211-e212.	1.2	6
78	Treatment of epidermal growth factor receptor inhibitorâ€induced severe paronychia with pyogenic granulomaâ€like lesions with topical betaxolol: an openâ€label observation study. International Journal of Dermatology, 2020, 59, 326-332.	1.0	6
79	Filaggrin: An Emerging Star in Atopic March. Journal of the Formosan Medical Association, 2008, 107, 429-431.	1.7	5
80	Lipodystrophia Centrifugalis Abdominalis Infantilis: Report of four cases. Pediatric Dermatology, 2012, 29, 308-310.	0.9	5
81	Recurrent Scedosporium apiospermum mycetoma successfully treated by surgical excision and voriconazole. Dermatologica Sinica, 2014, 32, 29-32.	0.5	5
82	Novel indel mutation of STS underlies a new phenotype of self-healing recessive X-linked ichthyosis. Journal of Dermatological Science, 2015, 79, 317-319.	1.9	5
83	Erythrokeratoderma Variabilis Caused by p.Gly45Glu in Connexin 31: Importance of the First Extracellular Loop Glycine Residue for Gap Junction Function. Acta Dermato-Venereologica, 2016, 96, 557-559.	1.3	5
84	Investigation of water bonding status of normal and psoriatic skin in vivo using diffuse reflectance spectroscopy. Scientific Reports, 2021, 11, 8901.	3.3	5
85	The influence of gender and smoking on hidradenitis suppurativa: A retrospective study of 161 patients in Taiwan. Dermatologica Sinica, 2021, 39, 125.	0.5	5
86	Disseminated superficial porokeratosis and disseminated superficial actinic porokeratosis: A case series of 39 patients. Dermatologica Sinica, 2020, 38, 221.	0.5	5
87	Central retinal vein occlusion and subsequent neovascular glaucoma after adalimumab treatment for psoriasis. Clinical and Experimental Dermatology, 2014, 39, 72-73.	1.3	4
88	Improved molecular diagnosis of the common recurrent intragenic deletion mutation in <scp><i>IKBKG</i></scp> in a <scp>F</scp> ilipino family with incontinentia pigmenti. Australasian Journal of Dermatology, 2016, 57, 150-153.	0.7	4
89	Systematised naevus sebaceus resulting from postâ€zygotic mutation in <scp><i>HRAS</i></scp> . Australasian Journal of Dermatology, 2017, 58, 58-60.	0.7	4
90	Pterygium and thinning of nails as an unusual manifestation in Clouston syndrome. Journal of Dermatology, 2019, 46, e329-e330.	1.2	4

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#	Article	IF	CITATIONS
91	Genetic Diagnosis of Rubinstein–Taybi Syndrome With Multiplex Ligation-Dependent Probe Amplification (MLPA) and Whole-Exome Sequencing (WES): Case Series With a Novel CREBBP Variant. Frontiers in Genetics, 2022, 13, 848879.	2.3	4
92	Investigating the clinical implication of corneometer and mexameter readings towards objective, efficient evaluation of psoriasis vulgaris severity. Scientific Reports, 2022, 12, 7469.	3.3	4
93	Lymphomatoid papulosis in association with mycosis fungoides: A clinical and histopathologic review of five Taiwanese cases. Dermatologica Sinica, 2014, 32, 75-81.	0.5	3
94	Insect biteâ€like reaction in association with chronic lymphocytic leukemia. International Journal of Dermatology, 2015, 54, 1191-1193.	1.0	3
95	Excimer lamp as an effective alternative treatment for severe alopecia areata. Dermatologica Sinica, 2015, 33, 151-153.	0.5	3
96	Ectodermal dysplasia–skin fragility syndrome resulting from a new atypical homozygous cryptic acceptor splice site mutation in PKP1. Journal of Dermatological Science, 2016, 84, 210-212.	1.9	3
97	Perifolliculitis capitis abscedens et suffodiens: eine Fallserie mit 66 Patienten und ein Vorschlag zur Klassifikation. JDDG - Journal of the German Society of Dermatology, 2018, 16, 1219-1227.	0.8	3
98	Transplantation of autologous single hair units heals chronic wounds in autosomal recessive dystrophic epidermolysis bullosa: A proof-of-concept study. Journal of Tissue Viability, 2021, 30, 36-41.	2.0	3
99	Plasma metabolomic and lipidomic profiling highlights metabolic changes in keloidâ€prone individuals. Experimental Dermatology, 2022, 31, 433-434.	2.9	3
100	Novel p.Ala675Thr missense mutation in <i>TRPV3</i> in Olmsted syndrome. Clinical and Experimental Dermatology, 2020, 45, 796-798.	1.3	2
101	Complexity of Transcriptional and Translational Interference of Laminin-332 Subunits in Junctional Epidermolysis Bullosa with LAMB3 Mutations. Acta Dermato-Venereologica, 2021, 101, adv00522.	1.3	2
102	Topical ivermectinâ€induced transient flare of rosacea as a host reaction to killed Demodex mites preventable by shortâ€ŧerm use of topical corticosteroid. Dermatologic Therapy, 2022, 35, e15517.	1.7	2
103	Bowen's disease with features resembling myrmecia wart. Journal of Dermatology, 2015, 42, 90-93.	1.2	1
104	Conjunctival geographic ulcer: An overlooked sign of herpes simplex virus infection. Journal of Clinical Virology, 2015, 64, 40-44.	3.1	1
105	Blaschkoâ€linear lichen planus: Clinicopathological and genetic analysis. Journal of Dermatology, 2020, 47, e384-e385.	1.2	1
106	Impact of narrowâ€band ultraviolet B phototherapy on remission and relapses of mycosis fungoides in patients with Fitzpatrick skin <scp>III</scp> – <scp>IV</scp> . Journal of the European Academy of Dermatology and Venereology, 2020, 34, e286-e288.	2.4	1
107	Cutaneous cytomegalovirus (CMV) infection in a patient with metastasized lung cancer. Clinical Microbiology and Infection, 2021, 27, 565-567.	6.0	1
108	Transcriptomic Analysis of Blaschko-Linear Psoriasis Reveals Shared and Distinct Features with Psoriasis Vulgaris. Journal of Investigative Dermatology, 2021, , .	0.7	1

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#	Article	IF	CITATIONS
109	Case of inherited epidermolysis bullosa simplex with <i>KLHL24</i> gene mutation in Japan. Journal of Dermatology, 2022, 49, .	1.2	1
110	Transcriptomic response of peripheral blood mononuclear cells to secukinumab in an 8â€yearâ€old boy with juvenile generalized pustular psoriasis. Journal of the European Academy of Dermatology and Venereology, 2022, 36, .	2.4	1
111	Novel compound heterozygous <i>ITGB4</i> mutations underlie lethal junctional epidermolysis bullosa with pyloric atresia and aplasia cutis congenita. Journal of Dermatology, 2022, 49, .	1.2	1
112	Incontinentia pigmenti in a male infant and a proposed diagnostic algorithm. Clinical and Experimental Dermatology, 2022, 47, 1366-1368.	1.3	1
113	Pyostomatitis vegetans following coronavirus disease 2019 vaccination in a patient with ulcerative colitis. Journal of Dermatology, 2022, 49, .	1.2	1
114	Plasma metabolomic profiling reflects the malnourished and chronic inflammatory state in recessive dystrophic epidermolysis bullosa. Journal of Dermatological Science, 2022, 107, 82-88.	1.9	1
115	Update on filaggrin mutations and atopic dermatitis. Expert Review of Dermatology, 2010, 5, 315-323.	0.3	0
116	Pruritic lichenoid lesions on the vulva of an 11-year-old girl. Dermatologica Sinica, 2011, 29, 111-112.	0.5	0
117	A large fungating verruciform xanthoma of the scrotum in association with arteriovenous malformation mimicking giant condyloma. Dermatologica Sinica, 2014, 32, 67-68.	0.5	0
118	Pigmented hidroacanthoma simplex of the scalp mimicking clonal seborrheic keratosis. Dermatologica Sinica, 2016, 34, 64-65.	0.5	0
119	Eosinophilic pustular folliculitis associated with Demodex overgrowth or demodicosis on the face – A report of five cases. Dermatologica Sinica, 2021, 39, 132.	0.5	0
120	A de novo COL17A1 splice-site mutation causing a 7-bp deletion in a Taiwanese patient with junctional epidermolysis bullosa. European Journal of Dermatology, 2021, 31, 267-269.	0.6	0
121	Serial punch excisions followed by second-intention healing for acne keloidalis papules. Journal of the American Academy of Dermatology, 2021, , .	1.2	0
122	Thymoma-associated graft-versus-host disease-like erythroderma: A harbinger of poor prognosis. Dermatologica Sinica, 2021, 39, 105.	0.5	0
123	Skin Stem Cells and Their Roles in Skin Regeneration and Disorders. , 2015, , 125-143.		0
124	Exertional plantar blistering as an easily overlooked clue for epidermolysis bullosa simplex. Dermatologica Sinica, 2019, 37, 170.	0.5	0
125	Genetics of Scars and Keloids. , 2020, , 47-53.		0
126	Bilateral auricular ossificans with stenosis of external ear canals and hearing loss caused by primary hyperparathyroidism: A case report. Dermatologica Sinica, 2020, 38, 248.	0.5	0

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#	Article	IF	CITATIONS
127	Melanocytic nevus with amyloid deposit – Report of three cases. Dermatologica Sinica, 2020, 38, 115.	0.5	0
128	Title is missing!. , 2020, 15, e0244620.		0
129	Title is missing!. , 2020, 15, e0244620.		0
130	Title is missing!. , 2020, 15, e0244620.		0
131	Title is missing!. , 2020, 15, e0244620.		0
132	Cas9â€guided haplotyping of three truncation variants in autosomal recessive disease. Human Mutation, 2022, , .	2.5	0
133	Immunohistochemical and molecular studies of resolved cutaneous Kaposi sarcoma in a kidney transplant recipient: A clinicopathological observation. Asian Journal of Surgery, 2022, , .	0.4	0
134	Novel compound heterozygous indel <scp><i>ZMPSTE24</i></scp> mutations in a Taiwanese male infant with restrictive dermopathy. Journal of Dermatology, 2022, 49, .	1.2	0
135	Autosomal dominant epidermolysis bullosa simplex exacerbated by hyperkeratotic scabies. Journal of Dermatology, 2022, 49, .	1.2	0