

Hajime Nakano

List of Publications by Year in descending order

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Version: 2024-02-01

12
papers

110
citations

1684188

5
h-index

1474206

9
g-index

12
all docs

12
docs citations

12
times ranked

148
citing authors

#	ARTICLE	IF	CITATIONS
1	Novel mutation of the ferrochelatase gene in a Japanese boy with erythropoietic protoporphyria. <i>Journal of Dermatology</i> , 2022, 49, .	1.2	1
2	A mild case of acute generalized exanthematous pustulosis caused by gemcitabine. <i>Journal of Dermatology</i> , 2022, 49, .	1.2	2
3	Autoantibodies to BPAG1e Trigger Experimental Bullous Pemphigoid in Mice. <i>Journal of Investigative Dermatology</i> , 2021, 141, 1167-1176.e3.	0.7	19
4	Novel <i>ATP2A2</i> nonsense mutation in a Japanese case with Darier's disease. <i>Journal of Dermatology</i> , 2021, 48, e149-e150.	1.2	0
5	Novel gross deletion mutation c.105_4042+498del in the <i>TNXB</i> gene in a Japanese woman with classical Ehlere Danlos syndrome: A case of uneventful pregnancy and delivery. <i>Journal of Dermatology</i> , 2021, 48, e227-e228.	1.2	2
6	The second Japanese case of porphyria cutanea tarda with a novel genetic mutation in <i>UROD</i> . <i>Journal of Dermatology</i> , 2021, 48, 1802-1803.	1.2	0
7	The first Japanese case of familial porphyria cutanea tarda diagnosed by a <i>UROD</i> mutation. <i>Journal of Dermatological Science</i> , 2019, 93, 65-67.	1.9	1
8	A novel missense mutation of the <i>STS</i> gene in two siblings with X-linked ichthyosis, complicated by short stature, bone density reduction, epilepsy, and cryptorchidism. <i>Clinical and Experimental Dermatology</i> , 2019, 44, 78-79.	1.3	4
9	Nagashima-type palmoplantar keratoderma and malignant melanoma in Japanese patients. <i>British Journal of Dermatology</i> , 2019, 180, 415-416.	1.5	7
10	Cutaneous collagenous vasculopathy induced by the vascular endothelial growth factor receptor inhibitor axitinib. <i>International Journal of Dermatology</i> , 2018, 57, e167-e169.	1.0	7
11	Papillon-Lefevre Syndrome and Malignant Melanoma. <i>Dermatology</i> , 2008, 217, 58-62.	2.1	42
12	Novel Ferrochelatase Mutations in Japanese Patients with Erythropoietic Protoporphyria: High Frequency of the Splice Site Modulator IVS3-48C Polymorphism in the Japanese Population. <i>Journal of Investigative Dermatology</i> , 2006, 126, 2717-2719.	0.7	25