Eijiro Akasaka

List of Publications by Year in descending order

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1937685 1588992 14 66 4 8 citations h-index g-index papers 14 14 14 81 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Pyoderma gangrenosum triggered by switching from adalimumab to secukinumab. Journal of Dermatology, 2019, 46, e108-e109.	1.2	19
2	Diversity of Mechanisms Underlying Latent TGF \hat{l}^2 Activation in Recessive Dystrophic Epidermolysis Bullosa. Journal of Investigative Dermatology, 2021, 141, 1450-1460.e9.	0.7	17
3	Primary Amelanotic Rhabdoid Melanoma: A Case Report with Review of the Literature. Case Reports in Dermatology, 2015, 7, 292-297.	0.8	6
4	Elderly-Onset Generalized Pustular Psoriasis without a Previous History of Psoriasis Vulgaris. Case Reports in Dermatology, 2015, 7, 187-193.	0.8	6
5	Abnormal keratinization and cutaneous inflammation in Mal de Meleda. Journal of Dermatology, 2020, 47, 554-558.	1.2	3
6	A 1,035-Subject Study Suggesting a History of Bone Fracture as a Possible Factor Associated with the Development of Anti-BP180 Autoantibodies. Journal of Investigative Dermatology, 2022, 142, 984-987.e3.	0.7	3
7	Diaminodiphenyl Sulfone-Induced Hemolytic Anemia and Alopecia in a Case of Linear IgA Bullous Dermatosis. Case Reports in Dermatology, 2015, 7, 183-186.	0.8	2
8	Successful treatment of lowâ€dose methotrexate in combination with systemic steroids for juvenile multiple and symmetrical circumscribed morphea. Journal of Dermatology, 2017, 44, e256-e257.	1.2	2
9	Pembrolizumab-Induced Lichen Planus on the Scalp of a Patient with Non-Small-Cell Lung Carcinoma. Case Reports in Dermatology, 2022, 13, 487-491.	0.8	2
10	Availability of mRNA Obtained from Peripheral Blood Mononuclear Cells for Testing Mutation Consequences in Dystrophic Epidermolysis Bullosa. International Journal of Molecular Sciences, 2021, 22, 13369.	4.1	2
11	A mild case of acute generalized exanthematous pustulosis caused by gemcitabine. Journal of Dermatology, 2022, 49, .	1.2	2
12	Novel mutation of the ferrochelatase gene in a Japanese boy with erythropoietic protoporphyria. Journal of Dermatology, 2022, 49, .	1.2	1
13	Two cases of the intermediate phenotype of recessive dystrophic epidermolysis bullosa harboring the novel COL7A1 mutation c.3570G>A. Journal of Dermatological Science, 2022, , .	1.9	1
14	Novel mutation c.263A> G in the $\langle i \rangle \langle scp \rangle ACVRL \langle scp \rangle 1 \langle i \rangle$ gene in a Japanese patient with hereditary hemorrhagic telangiectasia 2. Journal of Dermatology, 2019, 46, e22-e24.	1.2	0