

令和元年度 研究業績

雑誌

欧文

著者名	論文タイトル	誌名	巻：頁、出版年 (西暦)
Tanabe Y, Yamane M, Kato M, Teshima H, Kuribayashi M, Tatsukawa H, Takama H, <u>Akiyama M</u> , Hitomi K	Studies on differentiation-dependent expression and activity of distinct transglutaminases by specific substrate peptides using three-dimensional reconstituted epidermis.	FEBS J	286(13): 2536-2548, 2019
Murase Y, Takeichi T, <u>Akiyama M</u>	Aberrant CARD14 function might cause defective barrier formation.	J Allergy Clin Immunol	143(4): 1656-1657, 2019
Yoshihara N, Nakano H, <u>Sawamura D</u> , Kamata A, Matsuzaki H, Etoh T, <u>Ikeda S</u>	A case of junctional epidermolysis bullosa with pyloric atresia due to integrin $\beta 4$ gene mutations.	Dermatol Open J	4: 7-9, 2019
Ujiie H, Iwata H, <u>Yamagami J</u> , Nakama T, <u>Aoyama Y</u> , <u>Ikeda S</u> , Ishii N, Iwatsuki K, <u>Kurosawa M</u> , <u>Sawamura D</u> , Tanikawa A, Tsuruta D, Nishie W, Fujimoto W, <u>Amagai M</u> , <u>Shimizu H</u> , Committee for Guidelines for the Management of Pemphigoid Diseases (Including Epidermolysis Bullosa Acquisita)	Japanese guidelines for the management of pemphigoid (including epidermolysis bullosa acquisita).	J Dermatol	46: 1102-1135, 2019
<u>Kurosawa M</u> , Uehara R, Takagi A, <u>Aoyama Y</u> , Iwatsuki K, <u>Amagai M</u> , Nagai M, Nakamura Y, Inaba Y, Yokoyama K, <u>Ikeda S</u>	Results of nationwide epidemiologic survey of autosomal recessive congenital in Japan.	J Am Acad Dermatol	81(5): 1086-1092, 2019
Maruyama N, Takai T, Kamiyo S, Suchiva P, Ohba M, Takeshige T, Suzuki M, Hara M, Matsuno K, Harada S, Harada N, Nakae S, Sudo K, Okuno T, Yokomizo T, Ogawa H, Okumura K, <u>Ikeda S</u>	Cyclooxygenase inhibition in mice heightens adaptive- and innate-type responses against inhaled protease allergen and IL-33.	Allergy	74: 2237-2240, 2019

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Saito A, Nakamura Y, Tanaka R, Inoue S, Okiyama N, Ishitsuka Y, Maruyama H, Watanabe R, Yoshida K, <u>Ishiko A</u> , Fujimoto M, Shinkuma S, Fujisawa Y	Unusual Bone Lesions with Osteonecrosis Mimicking Bone Metastasis of Squamous Cell Carcinoma in Recessive Dystrophic Epidermolysis Bullosa.	Acta Derm Venereol	99(12): 1166-1169, 2019
<u>Kurosawa M</u> , Uehara R, Takagi A, <u>Aoyama Y</u> , Iwatsuki K, <u>Amagai M</u> , Nagai M, Nakamura Y, Inaba Y, Yokoyama K, <u>Ikeda S</u>	Results of a nationwide epidemiological survey of autosomal recessive congenital ichthyosis and ichthyosis syndromes in Japan.	J Am Acad Dermatol	46(12): 1102-1135, 2019
Korekawa A, Akasaka E, Rokunohe D, Fukui T, Kaneko T, <u>Sawamura D</u> , Ishikawa M, Yamamoto T, Nakano H	Nagashima-type palmoplantar keratoderma and malignant melanoma in Japanese patients.	Br J Dermatol	180(2): 415-416, 2019
Teramae A, Kobayashi Y, Kunimoto H, Nakajima K, <u>Suzuki T</u> , Tsuruta D, Fukai K	The Molecular Basis of Chemical Chaperone Therapy for Oculocutaneous Albinism Type 1A.	J Invest Dermatol	139(5): 1143-1149, 2019
Okamura K, Abe Y, Hayashi M, Saito T, Nagatani K, Tanoue T, Wataya-Kaneda M, Hozumi Y, <u>Suzuki T</u>	Impact of a 4-bp deletion variant (rs984225803) in the promoter region of SLC45A2 on color variation among a Japanese population.	J Dermatol	46(8): e295-e296, 2019
Tsutsumi R, Sugita K, Abe Y, Hozumi Y, <u>Suzuki T</u> , Yamada N, Yoshida Y, Yamamoto O	Leukoderma induced by rhododendrol is different from leukoderma of vitiligo in pathogenesis: A novel comparative morphological study.	J Cutan Pathol	46(2): 123-129, 2019
Okamura K, Hayashi M, Abe Y, Kono M, Nakajima K, <u>Aoyama Y</u> , Nishigori C, Ishimoto H, Ishimatsu Y, Nakajima M, Hozumi Y, <u>Suzuki T</u>	NGS-based targeted resequencing identified rare subtypes of albinism: Providing accurate molecular diagnosis for Japanese patients with albinism.	Pigm Cell Melanoma Res	32(6): 848-853, 2019

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Nao N, Sato K, Yamagishi J, Tahara M, Nakatsu Y, Seki F, Katoh H, Ohnuma A, Shirogane Y, Hayashi M, Suzuki T, Kikuta H, Nishimura H, Takeda M	Consensus and variations in cell line specificity among human metapneumovirus strains.	PLoS One	14(4) : 2019
Maekawa A, Arase N, <u>Tamai K</u> , Nomura T, Kiyohara E, Wataya- Kaneda M, Arase H, Katayama I, Fujimoto M	Case of epidermolytic ichthyosis with impairment of pulmonary function and exacerbated skin manifestations in a late middle-aged adult.	J Dermatol	46(12) : e480-e482, 2019
Ebens CL, McGrath JA, <u>Tamai K</u> , Hovnanian A, Wagner JE, Riddle MJ, Keene DR, DeFor TE, Tryon R, Chen M, Woodley DT, Hook K, Tolar J	Bone marrow transplant with post-transplant cyclophosphamide for recessive dystrophic epidermolysis bullosa expands the related donor pool and permits tolerance of nonhaematopoietic cellular grafts.	Br J Dermatol	181(6) : 1238-1246, 2019
Morita A, Kawakami Y, Kaji T, <u>Hirai Y</u> , Miyake T, Takahashi M, Yamasaki O, Sugiura K, Morizane S	Pediatric-onset annular pustular psoriasis in a patient with Down syndrome.	J Dermatol	46(10), e367-e368, 2019
Kamata A, Kurihara Y, Funakoshi T, Takahashi H, Kuroda K, Hachiya T, <u>Amagai M</u> , <u>Yamagami J</u>	Basement membrane zone IgE deposition is associated with bullous pemphigoid disease severity and treatment results.	Br J Dermatol	182(5), 1221-1227, 2020
Kakuta R, Kurihara Y, Yamagami J, Miyamoto J, Funakoshi T, Tanikawa A, <u>Amagai M</u>	Results of the guideline-based treatment for pemphigus: a single-centre experience with 84 cases.	J Eur Acad Dermatol	2020
Sugiyama S, Tanaka R, Hayashi H, Izumi K, Nishie W, <u>Aoyama Y</u>	Acquired hemophilia A in DPP4 inhibitor-induced bullous pemphigoid as an immune reconstitution syndrome.	Acta Dermato-Venereologica	in press
Murase Y, Takeichi T, Kawamoto A, Tanahashi K, Okuno Y, Takama H, Shimizu E, Ishikawa J, Ogi T, <u>Akiyama M</u>	Reduced stratum corneum acylceramides in autosomal recessive congenital ichthyosis with a NIPAL4 mutation.	J Dermatol Sci	97(1) : 50-56, 2020
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Minakawa S, Matsuzaki Y, Hashimoto T, Ishii N, Nishie W, Kayaba H, <u>Sawamura D</u>	Dipeptidyl peptidase-4 inhibitor-associated anti-laminin- γ 1 (p200) pemphigoid in a patient with psoriasis vulgaris.	J Dermatol	47(1): e25-e26, 2020
Nishikawa T, Okamura K, Moriyama M, Watanabe K, Ibusuki A, Sameshima S, Masamoto I, Yamazaki I, Tanita K, Kanekura T, Kanegane H, <u>Suzuki T, Kawano Y</u>	Novel AP3B1 compound heterozygous mutations in a Japanese patient with Hermansky-Pudlak syndrome type 2.	J Dermatol	47(2): 185-189, 2020
Mai S, Nishie W, Mai Y, Natsuga K, Nomura T, Suzuki S, Araki Y, <u>Suzuki T, Shimizu H</u>	Speckled lentiginous nevus in a patient with Hermansky-Pudlak syndrome type 1.	J Dermatol	47(1): e20-e21, 2020
Has C, Bauer JW, Bodemer C, Bolling M, Bruckner-Tuderman L, Diem A, Fine JD, Heagerty A, Hovnanian A, Marinkovich P, Martinez AE, McGrath JA, Moss C, Murrell DF, Palisson F, Schwieger-Briel A, Sprecher E, <u>Tamai K</u> , Uitto J, Woodley DT, Zambruno G, Mellerio JE	Consensus re-classification of inherited epidermolysis bullosa and other disorders with skin fragility.	Br J Dermatol	doi:10.1111/bjd.18921. Review. 2020
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Saeki H, <u>Terui T</u> , Morita A, Sano S, Imafuku S, Asahina A, Komine M, Etoh T, Igarashi A, Torii H, Abe M, Nakagawa H, Watanabe A, Yotsuyanagi H, Ohtsuki M	Japanese Guidance for Use of Biologics for Psoriasis (The 2019 Version).	J Dermatol	47(3), 201-222, 2020
<u>Hide M</u> , Fukunaga A, Maehara J, Eto K, Hao J, Vardi M, Nomoto Y	Efficacy, pharmacokinetics, and safety of icatibant for the treatment of Japanese patients with an acute attack of hereditary angioedema: A phase 3 open-label study.	Allergol Int	2019 in press