

雑誌

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
Kono M, Yokoyama N, Ogawa Y, Takama H, Sugiura K, Akiyama M.	Unilateral generalized linear porokeratosis with nail dystrophy.	J Dermatol	43(3)	286-7.	2016
Tanahashi K, Sugiura K, Sato T, Akiyama M.	Noteworthy clinical findings of harlequin ichthyosis: Digital autoamputation caused by cutaneous constriction bands in a case with novel ABCA12 mutations.	Br J Dermatol	174(3)	689-91.	2016
Takeichi T, Sugiura K, Tso S, Simpson MA, McGrath JA, Akiyama M.	Bi-allelic nonsense mutations in <i>ABHD5</i> underlie a mild phenotype of Dorfman-Chanarin syndrome.	J Dermatol Sci	81(2)	134-6.	2016
Ogawa M, Sugiura K, Yokota K, Muro Y, Akiyama M.	Anti-transcription intermediary factor 1- $\gamma$ antibody-positive clinically amyopathic dermatomyositis complicated by interstitial lung disease and breast cancer.	J Eur Acad Dermatol Venerol	30(2)	373-5.	2016
Nin-Asai R, Muro Y, Sekiya A, Sugiura K, Akiyama M.	Serum thymus and activation-regulated chemokine (TARC/CCL17) levels reflect the disease activity in a patient with bullous pemphigoid.	J Eur Acad Dermatol Venerol	30(2)	327-8.	2016
Shibata A, Sugiura K, Suzuki A, Ichiki T, Akiyama M.	Apparent homozygosity due to compound heterozygosity of one point mutation and an overlapping exon deletion mutation in <i>ABCA12</i> .	J Dermatol Sci	80(3)	196-202.	2015
Muro Y, Sugiura K, Akiyama M.	What are the "True" Pathogenic Anti-desmoglein Antibodies?	Acta Derm Venereol	95(7)	872-4.	2015
Sugiura K, Endo K, Akasaka T, Akiyama M.	Successful treatment with infliximab of sibling cases with generalized pustular psoriasis caused by deficiency of interleukin-36 receptor antagonist.	J Eur Acad Dermatol Venerol	29(10)	2054-6.	2015

Sugiura K, Akiyama M.	Lamellar ichthyosis caused by a previously unreported homozygous ALOXE3 mutation in East Asia.	Acta Derm Venereol	95(7)	858-9.	2015
Noda K, Sugiura K, Kono M, Akiyama M.	Porokeratotic eccrine ostial and dermal duct nevus with a somatic homozygous or monoallelic variant of connexin 26.	J Dermatol Sci	79(3)	317-9.	2015
Muro Y, Sugiura K, Nara M, Sakamoto I, Suzuki N, Akiyama M.	High incidence of cancer in anti-small ubiquitin-like modifier activating enzyme antibody-positive dermatomyositis.	Rheumatology (Oxford)	54(9)	1745-7.	2015
Tanahashi K, Sugiura K, Akiyama M.	Topical minoxidil improves congenital hypotrichosis caused by LIPH mutations.	Br J Dermatol	173(3)	865-6.	2015
Sugiura K, Nakasuka A, Kono H, Kono M, Akiyama M.	Impetigo herpetiformis with IL36RN mutations in a Chinese patient: A founder haplotype of c.115+6T>C in East Asia.	J Dermatol Sci	79(3)	319-20.	2015
Kono M, Suganuma M, Takama H, Zarzoso I, Saritha M, Bodet D, Aboobacker S, Kaliaperumal K, Suzuki T, Tomita Y, Sugiura K, Akiyama M.	Dowling-Degos disease with mutations in POFUT1 is clinicopathologically distinct from reticulate acropigmentation of Kitamura.	Br J Dermatol	173(2)	584-6.	2015
Miyake T, Umemura H, Doi Y, Kurosogabe J, Tsujikawa K, Hamada T, Sugiura K, Aoyama Y, Akiyama M, Iwatsuki K.	A case of annular pustular psoriasis with heterozygous IL36RN mutation.	Eur J Dermatol	25(4)	349-50.	2015
Sugiura K, Arima M, Matsunaga K, Akiyama M.	The novel GJB3 mutation p.Thr202Asn in the M4 transmembrane domain underlies erythrokeratoderma variabilis.	Br J Dermatol	173(1)	309-11.	2015

Aizu T, Matsui A, Takiyoshi N, Akasaka E, Kaneko T, Nakano H, Sugiura K, Akiyama M, Sawamura D.	Elderly-onset generalized pustular psoriasis without previous history of psoriasis vulgaris.	Case Rep Dermatol	7(2)	187-93.	2015
Ohno Y, Nakamichi S, Ohkuni A, Kamiyama N, Naoe A, Tsujimura H, Yokose U, Sugiura K, Ishikawa J, Akiyama M, Kihara A	Essential role of the cytochrome P450 CYP4F2 in the production of acylceramide, the key lipid for skin permeability barrier formation.	Proc Natl Acad Sci U S A	112(25)	7707-12.	2015
Nanbu Ai, Sugiura K, Kono M, Muro Y, Akiyama M.	Annular elastolytic giant cell granuloma successfully treated with minocycline hydrochloride.	Acta Derm Venereol	95(6)	756-7.	2015
Himoto T, Tanaka N, Saito A, Muro Y, Sugiura K, Tani J, Miyoshi H, Morishita A, Yoneyama H, Haba R, Masaki T.	Diversity of humoral responses to the centromere proteins among HCV-related chronic liver disease, PBC and AIH patients.	Clin Res Hepatol Gastroenterol	39(2)	222-9	2015
Shibata A, Tanahashi K, Sugiura K, Akiyama M.	<i>TRPS1</i> haploinsufficiency results in increased <i>SAT3</i> and <i>SOX9</i> mRNA expression in hair follicles in trichorhinophalangeal syndrome.	Acta Derm Venereol	95(5)	620-1.	2015