

Jun-Ichi Sakabe

List of Publications by Year in descending order

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58
papers

1,664
citations

331538

21
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39
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58
all docs

58
docs citations

58
times ranked

2700
citing authors

#	ARTICLE	IF	CITATIONS
1	In-vivo Imaging of CD8+ T cell-mediated Keratinocyte Apoptosis in Graft-Versus Host Disease-like Dermatitis in Involucrin-mOVA Mice. <i>Journal of Investigative Dermatology</i> , 2022, , .	0.3	0
2	Plasmacytoid dendritic cells as a possible key player to initiate alopecia areata in the C3H/HeJ mouse. <i>Allergology International</i> , 2020, 69, 121-131.	1.4	19
3	Suprabasin-null mice retain skin barrier function and show high contact hypersensitivity to nickel upon oral nickel loading. <i>Scientific Reports</i> , 2020, 10, 14559.	1.6	11
4	Tp63-expressing adult epithelial stem cells cross lineages boundaries revealing latent hairy skin competence. <i>Nature Communications</i> , 2020, 11, 5645.	5.8	9
5	Protective role of Galectin-7 for skin barrier impairment in atopic dermatitis. <i>Clinical and Experimental Allergy</i> , 2020, 50, 922-931.	1.4	9
6	Potential role of transforming growth factor- β 1/Smad signaling in secondary lymphedema after cancer surgery. <i>Cancer Science</i> , 2020, 111, 2620-2634.	1.7	16
7	Decreased expression of suprabasin induces aberrant differentiation and apoptosis of epidermal keratinocytes: Possible role for atopic dermatitis. <i>Journal of Dermatological Science</i> , 2019, 95, 107-112.	1.0	17
8	Palmar hyperlinearity in early childhood atopic dermatitis is associated with filaggrin mutation and sensitization to egg. <i>Pediatric Dermatology</i> , 2019, 36, 213-218.	0.5	12
9	The Vitamin D3 analogue calcipotriol suppresses CpG-activated TLR9-MyD88 signalling in murine plasmacytoid dendritic cells. <i>Clinical and Experimental Dermatology</i> , 2018, 43, 445-448.	0.6	6
10	Sensitive skin is highly frequent in extrinsic atopic dermatitis and correlates with disease severity markers but not necessarily with skin barrier impairment. <i>Journal of Dermatological Science</i> , 2018, 89, 33-39.	1.0	28
11	Clinical course of the first Japanese family with Marie Unna hereditary hypotrichosis: a follow-up report. <i>European Journal of Dermatology</i> , 2018, 28, 406-407.	0.3	0
12	Voriconazole-induced photocarcinogenesis is promoted by aryl hydrocarbon receptor-dependent COX-2 upregulation. <i>Scientific Reports</i> , 2018, 8, 5050.	1.6	22
13	Distinctive downmodulation of plasmacytoid dendritic cell functions by vitamin D3 analogue calcipotriol. <i>Journal of Dermatological Science</i> , 2016, 84, 71-79.	1.0	9
14	Platelets Regulate the Migration of Keratinocytes via Podoplanin/CLEC-2 Signaling during Cutaneous Wound Healing in Mice. <i>American Journal of Pathology</i> , 2016, 186, 101-108.	1.9	28
15	Identification and Characterization of a Recessive Missense Mutation p.P277L in SERPINB7 in Nagashima-Type Palmoplantar Keratosis. <i>Journal of Investigative Dermatology</i> , 2016, 136, 325-328.	0.3	12
16	Pustular psoriasis-like lesions associated with hereditary lactate dehydrogenase M subunit deficiency without interleukin-36 receptor antagonist mutation: long-term follow-up of two cases. <i>British Journal of Dermatology</i> , 2015, 172, 1674-1676.	1.4	7
17	Gross Cystic Disease Fluid Protein 15 in Stratum Corneum Is a Potential Marker of Decreased Eccrine Sweating for Atopic Dermatitis. <i>PLoS ONE</i> , 2015, 10, e0125082.	1.1	3
18	Antihistaminic drug olopatadine downmodulates T cell chemotaxis toward CCL17 in patients with atopic dermatitis. <i>Allergology International</i> , 2015, 64, 200-202.	1.4	1

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19	Melanocyte-specific cytotoxic T lymphocytes in patients with rhododendrol-induced leukoderma. <i>Journal of Dermatological Science</i> , 2015, 77, 190-192.	1.0	20
20	Leukoderma in patients with atopic dermatitis. <i>Journal of Dermatology</i> , 2015, 42, 215-218.	0.6	7
21	Decreased Expression of Acetylcholine Esterase in Cholinergic Urticaria with Hypohidrosis or Anhidrosis. <i>Journal of Investigative Dermatology</i> , 2014, 134, 276-279.	0.3	38
22	Calcipotriol Increases hCAP18 mRNA Expression but Inhibits Extracellular LL37 Peptide Production in IL-17/IL-22-stimulated Normal Human Epidermal Keratinocytes. <i>Acta Dermato-Venereologica</i> , 2014, 94, 512-516.	0.6	22
23	Successful differentiation of herpes zoster-associated erythema multiforme from generalized extension of herpes by rapid polymerase chain reaction analysis. <i>Journal of Dermatology</i> , 2014, 41, 542-544.	0.6	6
24	A Japanese case of Mal de Meleda with SLURP1 mutation. <i>Journal of Dermatology</i> , 2014, 41, 764-765.	0.6	4
25	Atopic dermatitis presenting as generalized poikiloderma with filaggrin gene mutation. <i>Journal of Dermatology</i> , 2014, 41, 230-231.	0.6	3
26	Proteome analysis of stratum corneum from atopic dermatitis patients by hybrid quadrupole-orbitrap mass spectrometer. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 957-960.e8.	1.5	40
27	VEGF-A promotes IL-17A-producing $\gamma\delta$ T cell accumulation in mouse skin and serves as a chemotactic factor for plasmacytoid dendritic cells. <i>Journal of Dermatological Science</i> , 2014, 74, 116-124.	1.0	30
28	Potential application of in vivo imaging of impaired lymphatic duct to evaluate the severity of pressure ulcer in mouse model. <i>Scientific Reports</i> , 2014, 4, 4173.	1.6	43
29	High frequencies of positive nickel/cobalt patch tests and high sweat nickel concentration in patients with intrinsic atopic dermatitis. <i>Journal of Dermatological Science</i> , 2013, 72, 240-245.	1.0	36
30	Trichorhinophalangeal syndrome with low expression of TRPS1 on epidermal and hair follicle epithelial cells. <i>Journal of Dermatology</i> , 2013, 40, 396-398.	0.6	17
31	D1-like dopamine receptors antagonist inhibits cutaneous immune reactions mediated by Th2 and mast cells. <i>Journal of Dermatological Science</i> , 2013, 71, 37-44.	1.0	41
32	Mutations in SERPINB7, Encoding a Member of the Serine Protease Inhibitor Superfamily, Cause Nagashima-type Palmoplantar Keratosis. <i>American Journal of Human Genetics</i> , 2013, 93, 945-956.	2.6	102
33	Acicular, but not globular, titanium dioxide nanoparticles stimulate keratinocytes to produce pro-inflammatory cytokines. <i>Journal of Dermatology</i> , 2013, 40, 357-362.	0.6	10
34	Identification of a novel heterozygous mutation in the first Japanese case of Ulnna hereditary hypotrichosis. <i>Journal of Dermatology</i> , 2013, 40, 278-280.	0.6	6
35	Second report of FLG R501X mutation in Japanese patients with atopic dermatitis. <i>Journal of Dermatology</i> , 2013, 40, 498-499.	0.6	3
36	Erythrokeratoderma variabilis: First Japanese case documenting GJB3 mutation. <i>Journal of Dermatology</i> , 2013, 40, 402-403.	0.6	3

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37	Kallikrein-related Peptidase 5 Functions in Proteolytic Processing of Profilaggrin in Cultured Human Keratinocytes. <i>Journal of Biological Chemistry</i> , 2013, 288, 17179-17189.	1.6	66
38	Prostaglandin E2 is critical for the development of niacin-deficiency-induced photosensitivity via ROS production. <i>Scientific Reports</i> , 2013, 3, 2973.	1.6	19
39	Topical Simvastatin Accelerates Wound Healing in Diabetes by Enhancing Angiogenesis and Lymphangiogenesis. <i>American Journal of Pathology</i> , 2012, 181, 2217-2224.	1.9	116
40	Identification of mutations in the prostaglandin transporter gene <i>SLCO2A1</i> and its phenotype-genotype correlation in Japanese patients with pachydermoperiostosis. <i>Journal of Dermatological Science</i> , 2012, 68, 36-44.	1.0	72
41	Photosensitive drug eruption induced by bicalutamide within the UVB action spectrum. <i>European Journal of Dermatology</i> , 2012, 22, 402-404.	0.3	8
42	A group of atopic dermatitis without IgE elevation or barrier impairment shows a high Th1 frequency: Possible immunological state of the intrinsic type. <i>Journal of Dermatological Science</i> , 2012, 67, 37-43.	1.0	88
43	Sebaceous carcinoma arising at a chronic candidiasis skin lesion of a patient with keratitis-ichthyosis-deafness (KID) syndrome. <i>British Journal of Dermatology</i> , 2012, 166, 222-224.	1.4	6
44	Connexin 26 (<i>CX26</i>) mutations in keratitis-ichthyosis-deafness syndrome presenting with squamous cell carcinoma. <i>Journal of Dermatology</i> , 2012, 39, 814-815.	0.6	11
45	Molecular mapping of lymph node metastases by real-time reverse transcription polymerase chain reaction in two melanoma patients. <i>Journal of Dermatology</i> , 2011, 38, 1202-1205.	0.6	0
46	Topical Cholecystokinin Depresses Itch-Associated Scratching Behavior in Mice. <i>Journal of Investigative Dermatology</i> , 2011, 131, 956-961.	0.3	10
47	The Mandatory Role of IL-10-Producing and OX40 Ligand-Expressing Mature Langerhans Cells in Local UVB-Induced Immunosuppression. <i>Journal of Immunology</i> , 2010, 184, 5670-5677.	0.4	43
48	Involvement of Wnt Signaling in Dermal Fibroblasts. <i>American Journal of Pathology</i> , 2010, 176, 721-732.	1.9	29
49	Flaky Tail Mouse Denotes Human Atopic Dermatitis in the Steady State and by Topical Application with <i>Dermatophagoides pteronyssinus</i> Extract. <i>American Journal of Pathology</i> , 2010, 176, 2385-2393.	1.9	122
50	FTY720 Regulates Bone Marrow Egress of Eosinophils and Modulates Late-Phase Skin Reaction in Mice. <i>American Journal of Pathology</i> , 2010, 177, 1881-1887.	1.9	40
51	A missense mutation in Exon 1 of the keratin 9 gene in a Japanese patient with rarer type hereditary palmoplantar keratoderma. <i>European Journal of Dermatology</i> , 2009, 19, 286-287.	0.3	1
52	Expression of toll-like receptor 2, NOD2 and dectin-1 and stimulatory effects of their ligands and histamine in normal human keratinocytes. <i>British Journal of Dermatology</i> , 2009, 160, 297-304.	1.4	62
53	Possible involvement of T lymphocytes in the pathogenesis of Nagashima-type keratosis palmoplantaris. <i>Clinical and Experimental Dermatology</i> , 2009, 34, e282-e284.	0.6	11
54	Impaired Initiation of Contact Hypersensitivity by FTY720. <i>Journal of Investigative Dermatology</i> , 2008, 128, 2833-2841.	0.3	17

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55	A functional polymorphism of the TNF- β gene that is associated with type 2 DM. Biochemical and Biophysical Research Communications, 2008, 369, 943-947.	1.0	22
56	A novel 12-base pair deletion mutation in exon 15 of the porphobilinogen deaminase gene in a Taiwanese patient with acute intermittent porphyria. Blood Cells, Molecules, and Diseases, 2008, 41, 202.	0.6	1
57	“Nagashima-Type” Keratosis as a Novel Entity in the Palmoplantar Keratoderma Category. Archives of Dermatology, 2008, 144, 375-9.	1.7	53
58	CXCL12-CXCR4 Engagement Is Required for Migration of Cutaneous Dendritic Cells. American Journal of Pathology, 2007, 171, 1249-1257.	1.9	227