Nobuo Kanazawa

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

109 papers	2,911 citations	23 h-index	53 g-index
122 ext. papers	3,387 ext. citations	3.7 avg, IF	4.89 L-index

#	Paper	IF	Citations
109	The conduit system transports soluble antigens from the afferent lymph to resident dendritic cells in the T cell area of the lymph node. *Immunity*, **2005**, 22, 19-29	32.3	583
108	Early-onset sarcoidosis and CARD15 mutations with constitutive nuclear factor-kappaB activation: common genetic etiology with Blau syndrome. *Blood*, **2005**, 105, 1195-7	2.2	368
107	Proteasome assembly defect due to a proteasome subunit beta type 8 (PSMB8) mutation causes the autoinflammatory disorder, Nakajo-Nishimura syndrome. *Proceedings of the National Academy of Sciences of the United States of America*, **2011**, 108, 14914-9	11.5	227
106	Role of the NOD2 genotype in the clinical phenotype of Blau syndrome and early-onset sarcoidosis. *Arthritis and Rheumatism*, **2009**, 60, 242-50		124
105	Dendritic cell immunoreceptors: C-type lectin receptors for pattern-recognition and signaling on antigen-presenting cells. *Journal of Dermatological Science*, **2007**, 45, 77-86	4.3	92
104	Fractalkine and macrophage-derived chemokine: T cell-attracting chemokines expressed in T cell area dendritic cells. *European Journal of Immunology*, **1999**, 29, 1925-32	6.1	91
103	Calumenin, a Ca2+-binding protein retained in the endoplasmic reticulum with a novel carboxyl-terminal sequence, HDEF. *Journal of Biological Chemistry*, **1997**, 272, 18232-9	5.4	88
102	Disease-associated mutations in CIAS1 induce cathepsin B-dependent rapid cell death of human THP-1 monocytic cells. *Blood*, **2007**, 109, 2903-11	2.2	80
101	A case of neonatal-onset autoinflammatory syndrome with a de novo PSMB9 mutation resembling Nakajo-Nishimura syndrome. *Pediatric Rheumatology*, **2015**, 13,	3.5	78
100	Cutaneous polyarteritis nodosa: revisiting its definition and diagnostic criteria. *Archives of Dermatological Research*, **2009**, 301, 117-21	3.3	72
99	Dendritic cell immunoactivating receptor, a novel C-type lectin immunoreceptor, acts as an activating receptor through association with Fc receptor gamma chain. *Journal of Biological Chemistry*, **2003**, 278, 32645-52	5.4	70
98	DCIR acts as an inhibitory receptor depending on its immunoreceptor tyrosine-based inhibitory motif. *Journal of Investigative Dermatology*, **2002**, 118, 261-6	4.3	70
97	Steady state migratory RelB+ langerin+ dermal dendritic cells mediate peripheral induction of antigen-specific CD4+ CD25+ Foxp3+ regulatory T cells. *European Journal of Immunology*, **2011**, 41, 1420-34	6.1	65
96	Presence of a sporadic case of systemic granulomatosis syndrome with a CARD15 mutation. *Journal of Investigative Dermatology*, **2004**, 122, 851-2	4.3	56
95	Novel IL36RN mutation in a Japanese case of early onset generalized pustular psoriasis. *Journal of Dermatology*, **2013**, 40, 749-51	1.6	55
94	Autoinflammatory syndromes with a dermatological perspective. *Journal of Dermatology*, **2007**, 34, 601-18	1.8	54
93	Signaling and immune regulatory role of the dendritic cell immunoreceptor (DCIR) family lectins: DCIR, DCAR, dectin-2 and BDCA-2. *Immunobiology*, **2004**, 209, 179-90	3.4	50

92	Onychomycosis: modern diagnostic and treatment approaches. *Wiener Medizinische Wochenschrift*, **2013**, 163, 1-12	2.9	46
91	Nakajo-Nishimura syndrome: an autoinflammatory disorder showing pernio-like rashes and progressive partial lipodystrophy. *Allergology International*, **2012**, 61, 197-206	4.4	42
90	Molecular cloning, characterization, and chromosomal localization of FKBP23, a novel FK506-binding protein with Ca2+-binding ability. *Genomics*, **1998**, 54, 89-98	4.3	32
89	Human calumenin gene (CALU): cDNA isolation and chromosomal mapping to 7q32. *Genomics*, **1998** , 49, 331-3	4.3	31
88	The cytosolic pattern-recognition receptor Nod2 and inflammatory granulomatous disorders. *Journal of Dermatological Science*, **2005**, 39, 71-80	4.3	29
87	Molecular cloning of human dectin-2. *Journal of Investigative Dermatology*, **2004**, 122, 1522-4	4.3	26
86	Guideline for the diagnosis and treatment of scabies in Japan (second edition). *Journal of Dermatology*, **2008**, 35, 378-93	1.6	23
85	Guideline for the diagnosis and treatment of scabies in Japan (third edition): Executive Committee of Guideline for the Diagnosis and Treatment of Scabies. *Journal of Dermatology*, **2017**, 44, 991-1014	1.6	22
84	Novel compound heterozygous DNA ligase IV mutations in an adolescent with a slowly-progressing radiosensitive-severe combined immunodeficiency. *Clinical Immunology*, **2015**, 160, 255-60	9	22
83	Pluripotent Stem Cell Model of Nakajo-Nishimura Syndrome Untangles Proinflammatory Pathways Mediated by Oxidative Stress. *Stem Cell Reports*, **2018**, 10, 1835-1850	8	22
82	Sarcoidosis and molecular mimicry--important etiopathogenetic aspects: current state and future directions. *Wiener Klinische Wochenschrift*, **2012**, 124, 227-38	2.3	20
81	A new infant case of Nakajo-Nishimura syndrome with a genetic mutation in the immunoproteasome subunit: an overlapping entity with JMP and CANDLE syndrome related to PSMB8 mutations. *Dermatology*, **2013**, 227, 26-30	4.4	20
80	Hydroxychloroquine administration for Japanese lupus erythematosus in Wakayama: a pilot study. *Journal of Dermatology*, **2012**, 39, 531-5	1.6	19
79	ISSAID/EMQN Best Practice Guidelines for the Genetic Diagnosis of Monogenic Autoinflammatory Diseases in the Next-Generation Sequencing Era. *Clinical Chemistry*, **2020**, 66, 525-536	5.5	17
78	Mast cells and histamine metabolism in skin lesions from MRL/MP-lpr/lpr mice. *Autoimmunity Reviews*, **2009**, 8, 495-9	13.6	15
77	Expression of bcl-6 protein in normal skin and epidermal neoplasms. *Pathology International*, **1997**, 47, 600-7	1.8	15
76	Induction of PDGF-B in TCA-treated epidermal keratinocytes. *Archives of Dermatological Research*, **2007**, 299, 433-40	3.3	14
75	Clinical characteristics and treatment of 50 cases of Blau syndrome in Japan confirmed by genetic analysis of the mutation. *Annals of the Rheumatic Diseases*, **2020**, 79, 1492-1499	2.4	12

74	Novel PSTPIP1 gene mutation in pyoderma gangrenosum, acne and suppurative hidradenitis syndrome. *Journal of Dermatology*, **2018**, 45, e213-e214	1.6	12
73	Ultrasonographic assessment reveals detailed distribution of synovial inflammation in Blau syndrome. *Arthritis Research and Therapy*, **2014**, 16, R89	5.7	12
72	Influence of chemical peeling on the skin stress response system. *Experimental Dermatology*, **2012**, 21 Suppl 1, 8-10	4	11
71	Rare hereditary autoinflammatory disorders: towards an understanding of critical in vivo inflammatory pathways. *Journal of Dermatological Science*, **2012**, 66, 183-9	4.3	11
70	The protective effects of ultraviolet A1 irradiation on spontaneous lupus erythematosus-like skin lesions in MRL/lpr mice. *Clinical and Developmental Immunology*, **2009**, 2009, 673952		11
69	HLA-DQ and RBFOX1 as susceptibility genes for an outbreak of hydrolyzed wheat allergy. *Journal of Allergy and Clinical Immunology*, **2019**, 144, 1354-1363	11.5	10
68	Effects of sarpogrelate hydrochloride on skin ulcers and quality of life in patients with systemic sclerosis. *Journal of Dermatology*, **2012**, 39, 536-40	1.6	9
67	Race differences in immunogenetic features and photosensitivity of cutaneous lupus erythematosus from the aspect of Japanese studies. *Annals of the New York Academy of Sciences*, **2009**, 1173, 552-6	6.5	8
66	A Case of Blau Syndrome with NOD2 E383K Mutation. *Pediatric Dermatology*, **2016**, 33, e385-e387	1.9	8
65	Multiple courses of pulse corticosteroid therapy for alopecia areata. *Journal of Dermatology*, **2016**, 43, 1075-7	1.6	8
64	Prognostic factors of daily blood examination for advanced melanoma patients treated with nivolumab. *BioScience Trends*, **2018**, 12, 412-418	9.9	8
63	Designation of Autoinflammatory Skin Manifestations With Specific Genetic Backgrounds. *Frontiers in Immunology*, **2020**, 11, 475	8.4	7
62	Atypical fibroxanthoma-a diagnosis of exclusion!. *Wiener Medizinische Wochenschrift*, **2013**, 163, 380-6	2.9	7
61	Identification of 736T>A mutation of lipase H in Japanese siblings with autosomal recessive woolly hair. *Journal of Dermatology*, **2011**, 38, 900-4	1.6	7
60	Case of anti-laminin-gamma1 pemphigoid associated with psoriatic erythroderma. *Journal of Dermatology*, **2010**, 37, 272-5	1.6	7
59	Distribution analysis of infantile hemangioma or capillary malformation on the head and face in Japanese patients. *Journal of Dermatology*, **2019**, 46, 849-852	1.6	6
58	Influence of trichloroacetic acid peeling on the skin stress response system. *Journal of Dermatology* , **2011**, 38, 740-7	1.6	6
57	Preliminary study of etidronate for prevention of corticosteroid-induced osteoporosis caused by oral glucocorticoid therapy. *Clinical and Experimental Dermatology*, **2011**, 36, 165-8	1.8	6

56	No involvement of non-synonymous TLR2 polymorphisms in Japanese leprosy patients. *Journal of Dermatological Science*, **2009**, 54, 48-9	4.3	6
55	Severe ulceration with impaired induction of growth factors and cytokines in keratinocytes after trichloroacetic acid application on TRPV1-deficient mice. *European Journal of Dermatology*, **2012**, 22, 614-21	0.8	6
54	Sarcoidosis and Autoinflammation. *Inflammation and Regeneration*, **2011**, 31, 66-71	10.9	6
53	Autoimmunity versus autoinflammation--friend or foe?. *Wiener Medizinische Wochenschrift*, **2014**, 164, 274-7	2.9	5
52	Intractable leg ulcers in Blau syndrome. *Journal of Dermatology*, **2016**, 43, 1096-7	1.6	5
51	The effect of hydroxychloroquine on lupus erythematosus-like skin lesions in MRL/lpr mice. *Modern Rheumatology*, **2016**, 26, 744-8	3.3	5
50	WNK1/HSN2 founder mutation in patients with hereditary sensory and autonomic neuropathy: A Japanese cohort study. *Clinical Genetics*, **2017**, 92, 659-663	4	4
49	Heterozygous missense variant of the proteasome subunit β type 9 causes neonatal-onset autoinflammation and immunodeficiency. *Nature Communications*, **2021**, 12, 6819	17.4	4
48	The Expression of Histamine Receptors in Skin Lesions of MRL/MPlpr/ lpr Mice. *Open Dermatology Journal*, **2008**, 2, 83-86	1.1	4
47	Depletion of Epidermal Langerhans Cells in the Skin Lesions of Pellagra Patients. *American Journal of Dermatopathology*, **2017**, 39, 428-432	0.9	3
46	Peripheral blood eosinophilia is associated with the presence of skin ulcers in patients with systemic sclerosis. *Journal of Dermatology*, **2019**, 46, 334-337	1.6	3
45	Change of serum cytokine profiles by propranolol treatment in patients with infantile hemangioma. *Drug Discoveries and Therapeutics*, **2020**, 14, 89-92	5	3
44	A Case of Hailey-Hailey Disease with a Novel Nonsense Mutation in the Gene. *Annals of Dermatology*, **2017**, 29, 642-644	0.4	3
43	Pediatric pustular psoriasis with Turner's syndrome. *Journal of Dermatology*, **2015**, 42, 1208-9	1.6	3
42	Involvement of FcεR1α immunopositive cells in alopecia areata with atopic dermatitis and a high titer of serum immunoglobulin E. *European Journal of Dermatology*, **2014**, 24, 500-3	0.8	3
41	Pluripotent stem cell-based screening identifies CUDC-907 as an effective compound for restoring the in vitro phenotype of Nakajo-Nishimura syndrome. *Stem Cells Translational Medicine*, **2021**, 10, 455-464	6.9	3
40	Monitoring Cellular Movement with Photoconvertible Fluorescent Protein and Single-Cell RNA Sequencing Reveals Cutaneous Group 2 Innate Lymphoid Cell Subtypes, Circulating ILC2 and Skin-Resident ILC2.. *JID Innovations*, **2021**, 1, 100035		3
39	Difference in distribution of malignant melanoma and melanocytic nevus in the palm and finger. *BioScience Trends*, **2019**, 13, 361-363	9.9	2

38	Induced pluripotent stem cells representing Nakajo-Nishimura syndrome. *Inflammation and Regeneration*, **2019**, 39, 11	10.9	2
37	Beneficial effect of methotrexate on a child case of Nakajo-Nishimura syndrome. *Journal of Dermatology*, **2019**, 46, e365-e367	1.6	2
36	Severer lupus erythematosus-like skin lesions in MRL/lpr mice with homozygous Kit mutation. *Modern Rheumatology*, **2018**, 28, 319-326	3.3	2
35	Efficacy of 0.1% adapalene in a non-inflammatory Kyoto Rhino Rat acne model. *Journal of Dermatological Science*, **2014**, 76, 143-8	4.3	2
34	Hereditary disorders presenting with urticaria. *Immunology and Allergy Clinics of North America*, **2014**, 34, 169-79	3.3	2
33	Monogenic early-onset sarcoidosis is no longer a variant of "idiopathic" sarcoidosis. *Journal of the American Academy of Dermatology*, **2013**, 69, 164-5	4.5	2
32	Effective treatment of intractable skin ulcers using allogeneic cultured dermal substitutes in patients with systemic lupus erythematosus. *European Journal of Dermatology*, **2009**, 19, 594-6	0.8	2
31	2 Mutation-Associated Case with Blau Syndrome Triggered by BCG Vaccination. *Children*, **2021**, 8,	2.8	2
30	Dupilumab Improves Pruritus in Netherton Syndrome: A Case Study.. *Children*, **2022**, 9,	2.8	2
29	Repeated hyperhidrosis and chilblain-like swelling with ulceration of the fingers and toes in hereditary sensory and autonomic neuropathy type II. *Journal of Dermatology*, **2018**, 45, e308-e309	1.6	1
28	Is CANDLE the best nomenclature?. *British Journal of Dermatology*, **2014**, 171, 659-60	4	1
27	Genotyping of Trichophyton tonsurans isolate from a Japanese boy reveals infection in the USA. *Clinical and Experimental Dermatology*, **2009**, 34, 630-2	1.8	1
26	Lupus erythematosus tumidus in Japan: a case report and a review of the literature. *Modern Rheumatology*, **2009**, 19, 567-572	3.3	1
25	C-Type Lectin Receptors **2016**, 255-274		1
24	Effectiveness and safety of tacrolimus ointment combined with dupilumab for patients with atopic dermatitis in real-world clinical practice. *Journal of Dermatology*, **2021**, 48, 1564-1568	1.6	1
23	Case of Legionella pneumophila pneumonia (legionellosis) developed in a psoriatic arthritis patient receiving adalimumab. *Journal of Dermatology*, **2017**, 44, 982-983	1.6	0
22	Identification of a novel CCDC22 mutation in a patient with severe Epstein-Barr virus-associated hemophagocytic lymphohistiocytosis and aggressive natural killer cell leukemia. *International Journal of Hematology*, **2019**, 109, 744-750	2.3	0
21	Psoriatic arthritis induced by anti-programmed death 1 antibody pembrolizumab. *Journal of Dermatology*, **2019**, 46, e466-e467	1.6	0

20	Anticomplement therapy in bullous pemphigoid. *British Journal of Dermatology*, **2019**, 181, 448-449	4	0
19	Japanese guidelines for the management of palmoplantar keratoderma. *Journal of Dermatology*, **2021**, 48, e353-e367	1.6	0
18	Granular C3 dermatosis-like immunological manifestation found in a case of acute generalized exanthematous pustulosis: Implication for the mechanism in C3 deposition to the epidermal basement membrane zone. *Journal of Dermatology*, **2021**, 48, e238-e239	1.6	0
17	Dupilumab in atopic dermatitis patients with chronic hepatitis B. *Journal of Cutaneous Immunology and Allergy*,	0.3	0
16	CO2 narcosis as a notable cause of premature death in Nakajo-Nishimura syndrome. *Modern Rheumatology Case Reports*, **2019**, 3, 74-78	0.4	
15	Dental treatment for patients with Nakajo-Nishimura syndrome: Report of three cases. *Journal of Oral and Maxillofacial Surgery, Medicine, and Pathology*, **2020**, 32, 129-131	0.4	
14	Autoinflammatory Diseases. *The Japanese Journal of Sarcoidosis and Other Granulomatous Disorders* , **2016**, 36, 21-26	0	
13	Pachyonychia congenita in Japan: report of familial cases with a recurrent KRT16 mutation. *European Journal of Dermatology*, **2014**, 24, 122-3	0.8	
12	A Japanese Family with Congenital Erythrocytosis Caused by Haemoglobin Bethesda. *Internal Medicine*, **2015**, 54, 2389-93	1.1	
11	No involvement of the NOD1 polymorphism Glu266Lys in Japanese leprosy patients. *Journal of Dermatological Science*, **2009**, 56, 72-3	4.3	
10	Some issues on cutaneous sarcoidosis. *The Japanese Journal of Sarcoidosis and Other Granulomatous Disorders*, **2021**, 41, 52-54	0	
9	Lupus erythematosus tumidus in Japan: a case report and a review of the literature. *Modern Rheumatology*, **2009**, 19, 567-72	3.3	
8	Early-onset sarcoidosis and NOD2: Summary on genetic analysis of Japanese 10 cases. *Ensho Saisei*, **2005**, 25, 169-172		
7	Autoinflammatory Disorders Showing Pernio-like Eruptions. *Nishinihon Journal of Dermatology*, **2018**, 80, 321-326	0	
6	Hereditary diseases that should be differentiated from urticaria. *Nihon Shoni Arerugi Gakkaishi the Japanese Journal of Pediatric Allergy and Clinical Immunology*, **2020**, 34, 253-263	0.1	
5	Immunodeficiencies, autoinflammatory diseases and Hansen□ disease. *Japanese Journal of Leprosy*, **2017**, 86, 107-113	0	
4	Efficacy and Safety of Ebastine (EBASTEL) for Pruritic Skin Disease-Evaluation of High-dose Administration-. *Nishinihon Journal of Dermatology*, **2009**, 71, 609-615	0	
3	Case of Muckle-Wells syndrome with obesity. *Journal of Dermatology*, **2021**, 48, e438-e439	1.6	

2	Anaphylaxis to ginger induced by herbal medicine. *Allergology International*, **2020**, 69, 159-160	4.4
1	Evaluation of long-term disease control with dupilumab therapy using the Atopic Dermatitis Control Tool in real-world clinical practice. *Journal of Cutaneous Immunology and Allergy*,	0.3