Toshifumi Nomura

List of Publications by Year
in descending order

162	3,186	159585	182427
papers	citations	30 h-index	51 g-index

164	164	164	3654
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Hidradenitis suppurativa successfully treated with twoâ€stage surgery under disease control with adalimumab. Journal of Dermatology, 2022, 49, .	1.2	2
2	Intertwined vascular skin manifestations in a patient with SjÃ¶gren syndrome: A case report. Journal of Cutaneous Immunology and Allergy, 2022, 5, 22-23.	0.3	1
3	Case of antiâ€nuclear matrix protein 2 antibodyâ€positive juvenile dermatomyositis preceded by linear cutaneous lupus erythematosus on the face. Journal of Dermatology, 2022, 49, e18.	1.2	0
4	Multiple skin nodules on fingers in systemic lupus erythematosus. Rheumatology, 2022, , .	1.9	0
5	Pencil-core granuloma. Cmaj, 2022, 194, E14-E14.	2.0	1
6	Cutaneous arteritis following mRNAâ€1273 Moderna COVIDâ€19 vaccination. Journal of the European Academy of Dermatology and Venereology, 2022, 36, .	2.4	2
7	Development of systemic lupus erythematosus after dupilumab treatment in a case of atopic dermatitis. Journal of Dermatology, 2022, 49, 556-559.	1.2	3
8	Response to â€˜Clinical and direct immunofluorescence characteristics of cutaneous toxicity associated with enfortumab vedotinâ€™. British Journal of Dermatology, 2022, , .	1.5	1
9	Reliability of antinuclear matrix protein 2 antibody assays in idiopathic inflammatory myopathies is dependent on target protein properties. Journal of Dermatology, 2022, 49, 441-447.	1.2	3
10	Linear lichen planus in the lines of Blaschko suggestive of immuneâ€related adverse event. Journal of Cutaneous Immunology and Allergy, 2022, 5, 109-111.	0.3	2
11	Intravenous Injection of Muse Cells as a Potential Therapeutic Approach for Epidermolysis Bullosa. Journal of Investigative Dermatology, 2021, 141, 198-202.e6.	0.7	13
12	Hereditary Mucoepithelial Dysplasia and Autosomal-Dominant IFAP Syndrome Is a Clinical Spectrum Due to SREBF1 Variants. Journal of Investigative Dermatology, 2021, 141, 1596-1598.	0.7	6
13	KEAP1 and epidermal differentiation: Psoriatic epidermis as a model. Journal of Cutaneous Immunology and Allergy, 2021, 4, 132-134.	0.3	2
14	A case of anti-PL-7 antibody-positive anti-synthetase syndrome with dermatomyositis-associated erythema induced sclerodermatous changes. Rheumatology, 2021, 60, e362-e364.	1.9	0
15	Altered nucleocytoplasmic staining patterns of p62/SQSTM1 in cutaneous squamous cell carcinoma precursors. Journal of Cutaneous Immunology and Allergy, 2021, 4, 137-138.	0.3	0
16	Altered replication stress response due to CARD14 mutations promotes recombination-induced revertant mosaicism. American Journal of Human Genetics, 2021, 108, 1026-1039.	6.2	4
17	Japanese guidelines for the management of palmoplantar keratoderma. Journal of Dermatology, 2021, 48, e353-e367.	1.2	4
18	Primary cutaneous aspergillosis caused by Aspergillus welwitschiae: A case report. Journal of Dermatology, 2021, 48, e554-e555.	1.2	1

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19	Successful treatment of pulmonary hypertension with immunosuppressive therapy in a case of antiâ€synthetase syndrome. Journal of Dermatology, 2021, 48, e545-e546.	1.2	2
20	AAGAB is an assembly chaperone regulating AP1 and AP2 clathrin adaptors. Journal of Cell Science, 2021, 134, .	2.0	10
21	Evaluation of apremilast, an oral phosphodiesterase 4 inhibitor, for refractory cutaneous dermatomyositis: A phase 1b clinical trial. Journal of Dermatology, 2021, , .	1.2	4
22	Symmetrical acral keratoderma: A waxing and waning scaly pigmented skin lesions on the acral extremities. Journal of Dermatology, 2021, 48, e151-e152.	1.2	1
23	Development of Verruca Plana from Human Papillomavirus 78 Dependent on Host Immune State. Acta Dermato-Venereologica, 2021, 101, adv00608.	1.3	1
24	Loricrin Protects against Chemical Carcinogenesis. Journal of Investigative Dermatology, 2021, , .	0.7	2
25	Speckled lentiginous nevus in a patient with Hermanskyâ€“Pudlak syndrome type 1. Journal of Dermatology, 2020, 47, e20-e21.	1.2	1
26	Two Cases of Interleukin-7â€“Deficient Generalized Verrucosis. Clinical Infectious Diseases, 2020, 71, 1561-1563.	5.8	7
27	Refractory juvenile psoriatic uveitis without arthritis: a literature review. Journal of the European Academy of Dermatology and Venereology, 2020, 34, e173-e175.	2.4	0
28	Case of Xâ€linked hypohidrotic ectodermal dysplasia, along with facial bilateral reticular pigmentation. Journal of Cutaneous Immunology and Allergy, 2020, 3, 41-42.	0.3	0
29	Acute skin barrier disruption alters the secretion of lamellar bodies via the multilayered expression of ABCA12. Journal of Dermatological Science, 2020, 100, 50-57.	1.9	0
30	First case of symmetrical acral keratoderma in Japan with filaggrin mutation who showed marked improvement in skin manifestations using moisturizer. Journal of Dermatology, 2020, 47, e291-e293.	1.2	1
31	Hidradenitis Suppurativa as a Potential Subtype of Autoinflammatory Keratinization Disease. Frontiers in Immunology, 2020, 11, 847.	4.8	40
32	Recombination-induced revertant mosaicism in ichthyosis with confetti and loricrin keratoderma. Journal of Dermatological Science, 2020, 97, 94-100.	1.9	12
33	Cultured Epidermal Autografts from Clinically Revertant Skin as a Potential Wound Treatment for Recessive Dystrophic Epidermolysis Bullosa. Journal of Investigative Dermatology, 2019, 139, 2115-2124.e11.	0.7	22
34	AAGAB Controls AP2 Adaptor Assembly in Clathrin-Mediated Endocytosis. Developmental Cell, 2019, 50, 436-446.e5.	7.0	39
35	Case of epidermolytic ichthyosis with impairment of pulmonary function and exacerbated skin manifestations in a late middleâ€aged adult. Journal of Dermatology, 2019, 46, e480-e482.	1.2	0
36	Efficient Gene Reframing Therapy for Recessive Dystrophic Epidermolysis Bullosa with CRISPR/Cas9. Journal of Investigative Dermatology, 2019, 139, 1711-1721.e4.	0.7	39

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37	The identification of autoantigens in mucous membrane pemphigoid using immortalized oral mucosal keratinocytes. Journal of Oral Pathology and Medicine, 2019, 48, 60-67.	2.7	10
38	Lossâ€ofâ€function mutation in DSG1 underlies focal palmoplantar keratoderma. Journal of the European Academy of Dermatology and Venereology, 2019, 33, e137-e138.	2.4	5
39	Somatic recombination underlies frequent revertant mosaicism in loricrin keratoderma. Life Science Alliance, 2019, 2, e201800284.	2.8	13
40	Severe Septic Vasculitis Preceding Thoracic Empyema: Staphylococcus aureus Enterotoxin Deposition in Vessel Walls as a Possible Pathomechanism. Acta Dermato-Venereologica, 2019, 99, 464-465.	1.3	1
41	Novel <scp>COL</scp>7A1 mutation in a family with bullous dermolysis of the newborn: Phenotypic variability associated with a <scp>COL</scp>7A1 mutation within the same family. Journal of Dermatology, 2018, 45, e260-e261.	1.2	5
42	Detection of anti-BP180 NC16A autoantibodies after the onset of dipeptidyl peptidase-IV inhibitor-associated bullous pemphigoid: a report of three patients. British Journal of Dermatology, 2018, 179, 790-791.	1.5	28
43	Disseminated fusariosis emerged from prolonged local genital infection after cord blood transplantation. Journal of Infection and Chemotherapy, 2018, 24, 660-663.	1.7	6
44	Filaggrin gene mutations may influence the persistence of food allergies in Japanese primary school children. British Journal of Dermatology, 2018, 179, 190-191.	1.5	7
45	A solitary reddish nodule on the lower leg. International Journal of Dermatology, 2018, 57, 276-277.	1.0	0
46	Dermoscopic features of Bednar tumor: Report of a case. Journal of Dermatology, 2018, 45, e179-e180.	1.2	2
47	Gentamicin-Induced Readthrough and Nonsense-Mediated mRNA Decay of SERPINB7 Nonsense Mutant Transcripts. Journal of Investigative Dermatology, 2018, 138, 836-843.	0.7	33
48	Establishment of integration-free induced pluripotent stem cells from human recessive dystrophic epidermolysis bullosa keratinocytes. Journal of Dermatological Science, 2018, 89, 263-271.	1.9	14
49	Appearance of antidesmocollin 1 autoantibodies leading to a vegetative lesion in a patient with pemphigus vulgaris. British Journal of Dermatology, 2018, 178, 294-295.	1.5	4
50	Compound heterozygous missense mutations p.Leu207Pro and p.Tyr544Cys in TGM1 cause a severe form of lamellar ichthyosis. Journal of Dermatology, 2018, 45, 1463-1467.	1.2	12
51	Disseminated Erythematous Papules and Pustules: A Quiz. Acta Dermato-Venereologica, 2018, 99, 123-124.	1.3	0
52	Association of filaggrin gene mutations and childhood eczema and wheeze with phthalates and phosphorus flame retardants in house dust: The Hokkaido study on Environment and Children's Health. Environment International, 2018, 121, 102-110.	10.0	33
53	Apocrine mixed tumour on the abdomen: an atypical location. European Journal of Dermatology, 2018, 28, 234-235.	0.6	0
54	Diagnostic features of acquired dermal melanocytosis of the face and extremities. Clinical and Experimental Dermatology, 2018, 43, 806-809.	1.3	3

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55	The development of induced pluripotent stem cell-derived mesenchymal stem/stromal cells from normal human and RDEB epidermal keratinocytes. Journal of Dermatological Science, 2018, 91, 301-310.	1.9	32
56	Chromosomal inversions as a hidden disease-modifying factor for somatic recombination phenotypes. JCI Insight, 2018, 3, .	5.0	9
57	Rapid remission of severe pain from livedoid vasculopathy by apixaban. Journal of the European Academy of Dermatology and Venereology, 2017, 31, e45-e46.	2.4	12
58	Hypertrophic lupus erythematosus successfully treated with hydroxychloroquine. Journal of Dermatology, 2017, 44, e48-e49.	1.2	2
59	Thymomaâ€associated multiâ€organ autoimmunity: two cases and a review of the literature. Journal of the European Academy of Dermatology and Venereology, 2017, 31, e324-e326.	2.4	8
60	Altered balance of epidermis-related chemokines in epidermolysis bullosa. Journal of Dermatological Science, 2017, 86, 37-45.	1.9	9
61	Lateâ€onset skin involvement on the forehead in multicentric Castleman disease. International Journal of Dermatology, 2017, 56, e152-e153.	1.0	2
62	A case of recessive dystrophic epidermolysis bullosa with a novel c.6885_6898del14 mutation in the COL7A1 gene. Journal of Dermatological Science, 2017, 88, 139-141.	1.9	2
63	Identification of <scp>SERPINB</scp>7 mutations in Korean patients with Nagashimaâ€type palmoplantar keratosis. Journal of Dermatology, 2017, 44, 840-841.	1.2	8
64	Generalized Pustular Psoriasis. Journal of Pediatrics, 2017, 188, 305-305.e1.	1.8	2
65	Usefulness of dermoscopy in distinguishing benign lesions from angiosarcoma. Clinical and Experimental Dermatology, 2017, 42, 676-678.	1.3	1
66	Eosinophilic annular erythema is clinically characterized by central pigmentation reflecting basal melanosis: a clinicopathological study of 10 cases. Journal of the European Academy of Dermatology and Venereology, 2017, 31, 1916-1923.	2.4	17
67	Deficient stratum corneum intercellular lipid in a Japanese patient with lamellar ichthyosis with a homozygous deletion mutation in <scp>SDR</scp> 9C7 . British Journal of Dermatology, 2017, 177, e62-e64.	1.5	23
68	The first familial cases of epidermolysis bullosa simplex, generalized severe with p.Asn176Ser in KRT5 revealing the clinical chronology. Journal of the European Academy of Dermatology and Venereology, 2017, 31, e251-e253.	2.4	0
69	Pityriasis Rubra Pilaris Type V as an Autoinflammatory Disease by CARD14 Mutations. JAMA Dermatology, 2017, 153, 66.	4.1	64
70	A severe case of Xâ€linked ichthyosis showing palmar hyperlinearity without FLG mutations. Journal of the European Academy of Dermatology and Venereology, 2017, 31, e119-e120.	2.4	3
71	Compound heterozygotes for filaggrin gene mutations do not always show severe atopic dermatitis. Journal of the European Academy of Dermatology and Venereology, 2017, 31, 158-162.	2.4	13
72	Alopecia Induced by Timolol Eye-drops. Acta Dermato-Venereologica, 2017, 97, 295-296.	1.3	8

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73	Filaggrin Mutation in Korean Patients with Atopic Dermatitis. Yonsei Medical Journal, 2017, 58, 395.	2.2	25
74	Psoriasiform mycosis fungoides masquerading as tumourous plaques. European Journal of Dermatology, 2017, 27, 295-296.	0.6	4
75	Complete remission of angiolymphoid hyperplasia with eosinophilia using topical tacrolimus. European Journal of Dermatology, 2017, 27, 194-196.	0.6	0
76	Subcutaneous Nodule on the Right Palm of a Young Boy: A Quiz. Acta Dermato-Venereologica, 2017, 97, 1150-1151.	1.3	0
77	Erythrokeratoderma Variabilis Caused by p.Gly45Glu in Connexin 31: Importance of the First Extracellular Loop Glycine Residue for Gap Junction Function. Acta Dermato-Venereologica, 2016, 96, 557-559.	1.3	5
78	Possible cases of Nagashima-type palmoplantar keratosis? Comment on the article by Guoet al.. Clinical and Experimental Dermatology, 2016, 41, 320-320.	1.3	2
79	Nodular morphoea: a first case associated with linear morphoea. European Journal of Dermatology, 2016, 26, 95-96.	0.6	1
80	Warts in toe webs associated with human papillomavirus type 7: a specific cutaneous manifestation of this type?. British Journal of Dermatology, 2016, 174, 678-681.	1.5	5
81	Genetic analysis of a novel splice-site mutation inTMC8reveals thein vivoimportance of the transmembrane channel-like domain ofTMC8. British Journal of Dermatology, 2016, 175, 803-806.	1.5	8
82	Massive petechiae as an initial symptom of WaldenstrÃ¶m's macroglobulinemia. International Journal of Dermatology, 2016, 55, e361-2.	1.0	0
83	Safety of ustekinumab for the treatment of psoriasis vulgaris with myotonic dystrophy. European Journal of Dermatology, 2016, 26, 187-188.	0.6	5
84	Silicone medical adhesive removers for hyperkeratosis in epidermolysis bullosa. European Journal of Dermatology, 2016, 26, 501-502.	0.6	0
85	RNA recognition motif of LEMD3 as a key player in the pathogenesis of Buschkeâ€“Ollendorff syndrome. Journal of Dermatological Science, 2016, 81, 205-208.	1.9	5
86	Revertant Mosaicism in Ichthyosis with Confetti Caused by a Frameshift Mutation inÂKRT1. Journal of Investigative Dermatology, 2016, 136, 2093-2095.	0.7	23
87	Tuberculoid reaction to a cosmetic tattoo on the lips. European Journal of Dermatology, 2015, 25, 485-487.	0.6	3
88	Striate palmoplantar keratoderma: Report of a novel DSG1 mutation and atypical clinical manifestations. Journal of Dermatological Science, 2015, 80, 223-225.	1.9	6
89	Non-solar-induced elastotic bands on the forearm. European Journal of Dermatology, 2015, 25, 508-509.	0.6	1
90	Dermoscopic observation in adenoma of the nipple. Journal of Dermatology, 2015, 42, 341-342.	1.2	10

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91	Identification of previously unknownSERPINB7 splice variants in patients with Nagashima-type palmoplantar keratosis reveals the importance of the CD-loop of SERPINB7. British Journal of Dermatology, 2015, 173, 1288-1290.	1.5	6
92	Deep venous thrombosis in patients with BehÃ§et's disease. Journal of Dermatology, 2015, 42, 101-102.	1.2	0
93	Plasma cell cheilitis extending beyond vermillion border. Journal of Dermatology, 2015, 42, 935-936.	1.2	7
94	Lowâ€dose etretinate shows promise in management of punctate palmoplantar keratoderma type 1: Case report and review of the published work. Journal of Dermatology, 2015, 42, 889-892.	1.2	8
95	Pruritic Papules Following Lumbar Corset Use: A Quiz. Acta Dermato-Venereologica, 2015, 95, 763-766.	1.3	2
96	Punctate Palmoplantar Keratoderma Type 1: A Novel AAGAB Mutation and Efficacy of Etretinate. Acta Dermato-Venereologica, 2015, 95, 110-111.	1.3	14
97	Lossâ€ofâ€function mutations in the gene encoding filaggrin underlie a Japanese family with foodâ€dependent exerciseâ€induced anaphylaxis. Journal of the European Academy of Dermatology and Venereology, 2015, 29, 805-808.	2.4	8
98	Anti-MDA5 antibody-positive dermatomyositis with lethal progressive interstitial lung disease and advanced gastric cancer. European Journal of Dermatology, 2014, 24, 490-491.	0.6	11
99	A new filaggrin gene mutation in a Korean patient with ichthyosis vulgaris. European Journal of Dermatology, 2014, 24, 491-493.	0.6	2
100	Solitary Tumour on the Neck: A Quiz. Acta Dermato-Venereologica, 2014, 94, 619-622.	1.3	0
101	An annexin A1â€“FPR1 interaction contributes to necroptosis of keratinocytes in severe cutaneous adverse drug reactions. Science Translational Medicine, 2014, 6, 245ra95.	12.4	95
102	Repeated skin sampling and prolonged incubation period identified cutaneous M ycobacterium chelonae infection on the face in an immunocompetent man. British Journal of Dermatology, 2014, 170, 471-473.	1.5	2
103	Highly prevalent<scp>SERPINB7</scp>founder mutation causes pseudodominant inheritance pattern in Nagashimaâ€type palmoplantar keratosis. British Journal of Dermatology, 2014, 171, 847-853.	1.5	28
104	Recurrent Course and CD30 Expression of Atypical T Lymphocytes Distinguish Lymphomatoid Papulosis From Primary Cutaneous Aggressive Epidermotropic CD8+ Cytotoxic T-cell Lymphoma. Acta Dermato-Venereologica, 2014, 94, 613-614.	1.3	2
105	An analysis of the best method for evaluating anteversion of the acetabular component after total hip replacement on plain radiographs. Bone and Joint Journal, 2014, 96-B, 597-603.	4.4	56
106	Comprehensive screening for a complete set of Japaneseâ€populationâ€specific filaggrin gene mutations. Allergy: European Journal of Allergy and Clinical Immunology, 2014, 69, 537-540.	5.7	50
107	Distinct behavior of human Langerhans cells and inflammatory dendritic epidermal cells at tight junctions in patients with atopic dermatitis. Journal of Allergy and Clinical Immunology, 2014, 134, 856-864.	2.9	114
108	Tubular apocrine adenoma clinically and dermoscopically mimicking basal cell carcinoma. Journal of the American Academy of Dermatology, 2014, 71, e45-e46.	1.2	9

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109	A novel NCSTN mutation alone may be insufficient for the development of familial hidradenitis suppurativa. Journal of Dermatological Science, 2014, 74, 180-182.	1.9	28
110	Extensive Erythema and Hyperkeratosis on the Extremities and Lumbar Area as an Unusual Mani-festation of Nagashima-type Palmoplantar Keratosis. Acta Dermato-Venereologica, 2014, 96, 856-8.	1.3	2
111	Dermoscopy of pseudoxanthoma elasticum-like papillary dermal elastolysis. Journal of the American Academy of Dermatology, 2013, 69, e202-e203.	1.2	11
112	A novel splice site mutation inNCSTNunderlies a Japanese family with hidradenitis suppurativa. British Journal of Dermatology, 2013, 168, 206-209.	1.5	45
113	A Lack of Premature Termination Codon Read-Through Efficacy of PTC124 (Ataluren) in a Diverse Array of Reporter Assays. PLoS Biology, 2013, 11, e1001593.	5.6	118
114	Impaired Epidermal Permeability Barrier in Mice Lacking Elovl1, the Gene Responsible for Very-Long-Chain Fatty Acid Production. Molecular and Cellular Biology, 2013, 33, 2787-2796.	2.3	137
115	The Î²9 Loop Domain of PA-PLA1Î± Has a Crucial Role in Autosomal Recessive Woolly Hair/Hypotrichosis. Journal of Investigative Dermatology, 2012, 132, 2093-2095.	0.7	11
116	Haploinsufficiency for AAGAB causes clinically heterogeneous forms of punctate palmoplantar keratoderma. Nature Genetics, 2012, 44, 1272-1276.	21.4	78
117	Intraepidermal neutrophilic IgA pemphigus successfully treated with dapsone. European Journal of Dermatology, 2012, 22, 282-283.	0.6	10
118	Type VII Collagen Deficiency Causes Defective Tooth Enamel Formation due to Poor Differentiation of Ameloblasts. American Journal of Pathology, 2012, 181, 1659-1671.	3.8	19
119	A group of atopic dermatitis without IgE elevation or barrier impairment shows a high Th1 frequency: Possible immunological state of the intrinsic type. Journal of Dermatological Science, 2012, 67, 37-43.	1.9	88
120	AVULSION OF FLEXOR DIGITORUM PROFUNDUS SECONDARY TO RECURRENT ENCHONDROMA. Hand Surgery, 2011, 16, 205-206.	0.6	1
121	PrevalentLIPHfounder mutations lead to loss of P2Y5 activation ability of PA-PLA₁Î± in autosomal recessive hypotrichosis. Human Mutation, 2010, 31, n/a-n/a.	2.5	53
122	Immature platelet fraction measurement in patients with chronic liver disease: a convenient marker for evaluating cirrhotic change. International Journal of Laboratory Hematology, 2010, 32, 299-306.	1.3	16
123	Japanese-Specific Filaggrin Gene Mutations in Japanese Patients Suffering from Atopic Eczema and Asthma. Journal of Investigative Dermatology, 2010, 130, 2834-2836.	0.7	43
124	Chromosome 11q13.5 variant: No association with atopic eczema in the Japanese population. Journal of Dermatological Science, 2010, 59, 210-212.	1.9	2
125	Secondary syphilis mimicking warts in an HIV-positive patient. Sexually Transmitted Infections, 2009, 85, 484-484.	1.9	6
126	Analysis of Taiwanese ichthyosis vulgaris families further demonstrates differences inFLGmutations between European and Asian populations. British Journal of Dermatology, 2009, 161, 448-451.	1.5	49

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127	FLGmutation p.Lys4021X in the C-terminal imperfect filaggrin repeat in Japanese patients with atopic eczema. British Journal of Dermatology, 2009, 161, 1387-1390.	1.5	72
128	Clinical Severity Correlates with Impaired Barrier in Filaggrin-Related Eczema. Journal of Investigative Dermatology, 2009, 129, 682-689.	0.7	154
129	Prevalent and Rare Mutations in the Gene Encoding Filaggrin in Japanese Patients with Ichthyosis Vulgaris and Atopic Dermatitis. Journal of Investigative Dermatology, 2009, 129, 1302-1305.	0.7	43
130	Specific Filaggrin Mutations Cause Ichthyosis Vulgaris and Are Significantly Associated with Atopic Dermatitis in Japan. Journal of Investigative Dermatology, 2008, 128, 1436-1441.	0.7	128
131	Mild Recessive Bullous Congenital Ichthyosiform Erythroderma due to a Previously Unidentified Homozygous Keratin 10 Nonsense Mutation. Journal of Investigative Dermatology, 2008, 128, 1648-1652.	0.7	22
132	Papuloerythroderma of Ofuji associated with early gastric cancer. International Journal of Dermatology, 2008, 47, 590-591.	1.0	18
133	Amicrobial pustulosis associated with IgA nephropathy and SjÃ¶gren's syndrome. Journal of the American Academy of Dermatology, 2007, 57, 523-526.	1.2	25
134	Unique mutations in the filaggrin gene in Japanese patients with ichthyosis vulgaris andÂatopic dermatitis. Journal of Allergy and Clinical Immunology, 2007, 119, 434-440.	2.9	233
135	Epidermolysis bullosa acquisita associated with psoriasis vulgaris. Clinical and Experimental Dermatology, 2007, 32, 516-518.	1.3	23
136	Animation as a useful tool for assessing functional status in psoriatic arthritis. Journal of Dermatological Science, 2006, 44, 172-174.	1.9	0
137	Eccrine porocarcinoma and eccrine poroma arising in a scar. British Journal of Dermatology, 2004, 150, 1232-1233.	1.5	19
138	Rapid remission of severe pruritus from angiolymphoid hyperplasia with eosinophilia by pulsed dye laser therapy. Clinical and Experimental Dermatology, 2003, 28, 595-596.	1.3	19
139	Association of Infantile Cutaneous Haemangioma on the Face and Neck with Respiratory Distress in Infancy. Acta Dermato-Venereologica, 2003, 84, 72-73.	1.3	2
140	Reduced pre-movement facilitation of motor evoked potentials in spinocerebellar degeneration. Journal of the Neurological Sciences, 2001, 187, 41-47.	0.6	9
141	Nitric Oxide Donor NOR 3 Inhibits Ketogenesis from Oleate in Isolated Rat Hepatocytes by a Cyclic GMPâ€Independent Mechanism. Basic and Clinical Pharmacology and Toxicology, 1998, 82, 40-46.	0.0	5
142	Comparative Study on Germ Cell Mutation Induced by Urethane (Ethyl Carbamate) Gas and X-rays in Drosophila melanogaster. Japanese Journal of Cancer Research, 1997, 88, 461-467.	1.7	7
143	Distribution of nonprincipal neurons in the rat hippocampus, with special reference to their dorsoventral difference. Brain Research, 1997, 751, 64-80.	2.2	50
144	Laminar distribution of non-principal neurons in the rat hippocampus, with special reference to their compositional difference among layers. Brain Research, 1997, 764, 197-204.	2.2	44

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145	Induction of cancer, actinic keratosis, and specific p53 mutations by UVB light in human skin maintained in severe combined immunodeficient mice. Cancer Research, 1997, 57, 2081-4.	0.9	56
146	Cytogenotoxicities of sublimed urethane gas to the mouse embryo. Mutation Research - Genetic Toxicology Testing and Biomonitoring of Environmental Or Occupational Exposure, 1996, 369, 59-64.	1.2	7
147	Antigen receptor-mediated B cell death is blacked by signaling via CD72 or treatment with dextran sukfate and is defective in autoimmunity-prone mice. International Immunology, 1996, 8, 867-875.	4.0	62
148	Trigeminal neuralgia: Differentiation between intracranial mass lesions and ordinary vascular compression as causative lesions. Neurosurgical Review, 1994, 17, 51-57.	2.4	76
149	Effect of Histamine on the Blood-Tumor Barrier in Transplanted Rat Brain Tumors. , 1994, 60, 400-402.		6
150	Altered response to histamine in brain tumor vessels: the selective increase of regional cerebral blood flow in transplanted rat brain tumor. Journal of Neurosurgery, 1993, 79, 722-728.	1.6	17
151	Effects of prostacyclin analogue iloprost on the regional cerebral blood flow in transplanted rat brain tumour. Neurological Research, 1993, 15, 401-404.	1.3	7
152	Pharmacokinetics and pharmacodynamics of vecuronium bromide. Journal of Anesthesia, 1992, 6, 28-37.	1.7	1
153	Programmed Cell Death in Whole Body and Organ Systems by Low Dose Radiation. Journal of Radiation Research, 1992, 33, 109-123.	1.6	23
154	Nonâ€transformed, but not ras/mycâ€transformed, Serumâ€free Mouse Embryo Cells Recover from Growth Suppression by Azatyrosine. Japanese Journal of Cancer Research, 1992, 83, 851-858.	1.7	4
155	Paternal Exposure to Radiation and Offspring Cancer in Mice: Reanalysis and New Evidences. Journal of Radiation Research, 1991, 32, 64-72.	1.6	24
156	Multigeneration carcinogenesis. Radiation and Environmental Biophysics, 1991, 30, 201-203.	1.4	1
157	An examination of respiratory distress and chromosomal abnormalities in the offspring of male mice treated with ethylnitrosourea. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 1990, 229, 115-122.	1.0	11
158	SCID (severe combined immunodeficiency) mice as a new system to investigate metastasis of human tumors.. Journal of Radiation Research, 1990, 31, 288-292.	1.6	21
159	Suppression of developmental anomalies by maternal macrophages in mice.. Journal of Experimental Medicine, 1990, 172, 1325-1330.	8.5	45
160	Embryonic mutation as a possible cause of in utero carcinogenesis in mice revealed by postnatal treatment with 12-O-tetradecanoylphorbol-13-acetate. Cancer Research, 1990, 50, 2135-8.	0.9	14
161	Role of radiation-induced mutations in multigeneration carcinogenesis. Iarc (international Agency) Tj ETQq1 1 0.784314 rgBT /Overlock	0.4	1
162	X-ray- and chemically induced germ-line mutation causing phenotypical anomalies in mice. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 1988, 198, 309-320.	1.0	84