Peter R Hull

List of Publications by Year
in descending order

47 papers	2,366 citations	430874 18 h-index	243625 44 g-index
47 all docs	47 docs citations	47 times ranked	2798 citing authors

#	Article	IF	Citations
1	Comprehensive analysis of the gene encoding filaggrin uncovers prevalent and rare mutations in ichthyosis vulgaris and atopic eczema. Nature Genetics, 2007, 39, 650-654.	21.4	598
2	Loss-of-function variants in the filaggrin gene are a significant risk factor for peanut allergy. Journal of Allergy and Clinical Immunology, 2011, 127, 661-667.	2.9	424
3	Null Mutations in the Filaggrin Gene (FLG) Determine Major Susceptibility to Early-Onset Atopic Dermatitis that Persists into Adulthood. Journal of Investigative Dermatology, 2007, 127, 564-567.	0.7	298
4	Therapeutic siRNAs for dominant genetic skin disorders including pachyonychia congenita. Journal of Dermatological Science, 2008, 51, 151-157.	1.9	107
5	Isotretinoin Use and Subsequent Depression and Suicide. American Journal of Clinical Dermatology, 2003, 4, 493-505.	6.7	98
6	Isotretinoin Use in Acne: Prospective Evaluation of Adverse Events. Journal of Cutaneous Medicine and Surgery, 2000, 4, 66-70.	1.2	77
7	Management of acne: Canadian clinical practice guideline. Cmaj, 2016, 188, 118-126.	2.0	67
8	Acne, Depression, and Suicide. Dermatologic Clinics, 2005, 23, 665-674.	1.7	65
9	A Large Mutational Study in Pachyonychia Congenita. Journal of Investigative Dermatology, 2011, 131, 1018-1024.	0.7	60
10	Heterozygous Null Alleles in Filaggrin Contribute to Clinical Dry Skin in Young Adults and the Elderly. Journal of Investigative Dermatology, 2009, 129, 1042-1045.	0.7	58
11	Filaggrin gene mutation associations with peanut allergy persist despite variations in peanut allergy diagnostic criteria or asthma status. Journal of Allergy and Clinical Immunology, 2013, 132, 239-242.e7.	2.9	54
12	HLA-DQB1*02 and DQB1*06:03P are associated with peanut allergy. European Journal of Human Genetics, 2013, 21, 1181-1184.	2.8	43
13	Filaggrin Null Alleles Are Not Associated with Psoriasis. Journal of Investigative Dermatology, 2007, 127, 1878-1882.	0.7	41
14	The use of noninvasive imaging techniques in the diagnosis of melanoma: a prospective diagnostic accuracy study. Journal of the American Academy of Dermatology, 2021, 85, 353-359.	1.2	39
15	Localization of the Gene Causing Keratolytic Winter Erythema to Chromosome 8p22-p23, and Evidence for a Founder Effect in South African Afrikaans-Speakers. American Journal of Human Genetics, 1997, 61, 370-378.	6.2	38
16	Duplicated Enhancer Region Increases Expression of CTSB and Segregates with Keratolytic Winter Erythema in South African and Norwegian Families. American Journal of Human Genetics, 2017, 100, 737-750.	6.2	35
17	Breach Delivery: Increased Solute Uptake Points to a Defective Skin Barrier in Atopic Dermatitis. Journal of Investigative Dermatology, 2007, 127, 8-10.	0.7	33
18	Tattoo reactions as a sign of sarcoidosis: Figure 1:. Cmaj, 2012, 184, 432-432.	2.0	29

#	Article	IF	Citations
19	Recessive mutations in the gene encoding frizzled 6 cause twenty nail dystrophyâ€"Expanding the differential diagnosis for pachyonychia congenita. Journal of Dermatological Science, 2013, 70, 58-60.	1.9	22
20	Trends in Invasive Cutaneous Melanoma in Saskatchewan 1970â€"1999. Journal of Cutaneous Medicine and Surgery, 2003, 7, 433-442.	1.2	20
21	Immunotherapy for Cutaneous T-Cell Lymphoma: Current Landscape and Future Developments. Journal of Cutaneous Medicine and Surgery, 2019, 23, 537-544.	1.2	18
22	Filaggrin gene lossâ€ofâ€function mutations constitute a factor in patients with multiple contact allergies. Contact Dermatitis, 2019, 80, 354-358.	1.4	15
23	Trends in Invasive Cutaneous Melanoma in Saskatchewan 1970?1999. Journal of Cutaneous Medicine and Surgery, 2003, 7, 433-442.	1.2	13
24	Decreasing Rates of Neomycin Sensitization in Western Canada. Journal of Cutaneous Medicine and Surgery, 2016, 20, 446-452.	1.2	12
25	Major gene segregation of actinic prurigo among North American Indians in Saskatchewan. American Journal of Medical Genetics Part A, 2000, 92, 212-219.	2.4	11
26	Compliance with Self-Examination Surveillance in Patients with Melanoma and Atypical Moles: An Anonymous Questionnaire Study. Journal of Cutaneous Medicine and Surgery, 2011, 15, 97-102.	1.2	11
27	AAGAB Mutations in 18 Canadian Families With Punctate Palmoplantar Keratoderma and a Possible Link to Cancer. Journal of Cutaneous Medicine and Surgery, 2020, 24, 28-32.	1.2	9
28	Recurrent and Fixed Neutrophilic Dermatosis Associated With Dasatinib. Journal of Cutaneous Medicine and Surgery, 2018, 22, 621-623.	1.2	8
29	Exclusion of CTSB and FDFT1 as positional and functional candidate genes for keratolytic winter erythema (KWE). Journal of Dermatological Science, 2012, 65, 58-62.	1.9	6
30	The elusive gene for keratolytic winter erythema. South African Medical Journal, 2013, 103, 961.	0.6	6
31	Atrophying pityriasis versicolor. Cmaj, 2014, 186, 776-776.	2.0	6
32	PCQoL. Journal of Cutaneous Medicine and Surgery, 2015, 19, 57-65.	1.2	6
33	Photoprotection Knowledge and Behaviours Among Organ Transplant Recipients. Journal of Cutaneous Medicine and Surgery, 2017, 21, 217-220.	1.2	6
34	Extracorporeal photopheresis for the treatment of Crohn's disease. Transfusion and Apheresis Science, 2015, 52, 183-186.	1.0	5
35	Chronic Spontaneous Urticaria. Journal of Cutaneous Medicine and Surgery, 2016, 20, 241-243.	1.2	5
36	Dyshidrotic Bullous Pemphigoid: Case Report and Review of Literature. Journal of Cutaneous Medicine and Surgery, 2018, 22, 614-617.	1.2	5

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37	Hyposensitization and Desensitization in Allergic Contact Dermatitis. Dermatitis, 2012, 23, 148-152.	1.6	3
38	Primary Hyperoxaluria Type 1 (PH1) Presenting With End-Stage Kidney Disease and Cutaneous Manifestations in Adulthood: A Case Report. Canadian Journal of Kidney Health and Disease, 2021, 8, 205435812110589.	1.1	3
39	Birtâ€“Hoggâ€“DubÃ© syndrome: an inherited cause of spontaneous pneumothorax. Cmaj, 2011, 183, E601-E603.	2.0	2
40	Flow Cytometric Analysis of Regulatory T Cells During Hyposensitization of Acquired Allergic Contact Dermatitis. Dermatitis, 2014, 25, 60-65.	1.6	2
41	A Case of Lichen Sclerosus et Atrophicus With Distinct Erythematous Borders. Journal of Cutaneous Medicine and Surgery, 2015, 19, 600-603.	1.2	2
42	Two SMARCAD1 Variants Causing Basan Syndrome in a Canadian and a Dutch Family. JID Innovations, 2021, 1, 100022.	2.4	2
43	Disseminated Eczema Following Radiotherapy: A Case Report. Journal of Cutaneous Medicine and Surgery, 2018, 22, 353-355.	1.2	1
44	Acute onset of blisters in an infant with acrodermatitis enteropathica: A case report. SAGE Open Medical Case Reports, 2021, 9, 2050313X2098411.	0.3	1
45	Major gene segregation of actinic prurigo among North American Indians in Saskatchewan. American Journal of Medical Genetics Part A, 2000, 92, 212-219.	2.4	1
46	An Examination of Melanoma Detection and Characteristics at a Nova Scotia Tertiary Care Centre, From 2015-2019. Journal of Cutaneous Medicine and Surgery, 0, , 120347542211089.	1.2	1
47	Indirect Immunofluorescence for the Detection of Autoimmune Urticaria. Journal of Cutaneous Medicine and Surgery, 2016, 20, 113-117.	1.2	0