Eli Sprecher

List of Publications by Year
in descending order

231 papers	8,363 citations	53794 45 h-index	58581 82 g-index
241 all docs	241 docs citations	241 times ranked	8083 citing authors

#	Article	IF	Citations
1	Laryngeal Pemphigoid Evolution and Response to Treatment. Journal of Voice, 2023, 37, 471.e7-471.e14.	1.5	4
2	A stereoscopic optical system for objective quantification of the change in cumulative acne scar depth following various treatment interventions. Journal of Cosmetic Dermatology, 2022, 21, 2099-2105.	1.6	1
3	Vorinostat, a histone deacetylase inhibitor, as a potential novel treatment for psoriasis. Experimental Dermatology, 2022, 31, 567-576.	2.9	7
4	Neonatal inflammatory skin and bowel disease type 1 caused by a complex genetic defect and responsive to combined antiâ€tumour necrosis factorâ€Î± and interleukinâ€12/23 blockade. British Journal of Dermatology, 2022, 186, 1026-1029.	1.5	4
5	Association of a Third Dose of BNT162b2 Vaccine With Incidence of SARS-CoV-2 Infection Among Health Care Workers in Israel. JAMA - Journal of the American Medical Association, 2022, 327, 341.	7.4	76
6	Translational implications of Th17-skewed inflammation due to genetic deficiency of a cadherin stress sensor. Journal of Clinical Investigation, 2022, 132, .	8.2	24
7	Loss-of-function variants in KLF4 underlie autosomal dominant palmoplantar keratoderma. Genetics in Medicine, 2022, 24, 1085-1095.	2.4	3
8	The effect of a third-dose BNT162b2 vaccine on anti-SARS-CoV-2 antibody levels in immunosuppressed patients. Clinical Microbiology and Infection, 2022, 28, 735.e5-735.e8.	6.0	12
9	Superimposed type 2 segmental atopic dermatitis: a case series and review of the literature. Clinical and Experimental Dermatology, 2022, , .	1.3	0
10	Up-regulation of ST18 in pemphigus vulgaris drives a self-amplifying p53-dependent pathomechanism resulting in decreased desmoglein 3 expression. Scientific Reports, 2022, 12, 5958.	3.3	1
11	Acute Respiratory Distress Syndrome in a Carrier of an Interleukin-36 Receptor Antagonist Mutation With Generalized Pustular Psoriasis. Journal of Psoriasis and Psoriatic Arthritis, 2022, 7, 9-12.	0.7	0
12	Clinical efficacy of fecal microbial transplantation treatment in adults with moderateâ€toâ€severe atopic dermatitis. Immunity, Inflammation and Disease, 2022, 10, .	2.7	28
13	Short-Term Safety of Booster Immunization With BNT162b2 mRNA COVID-19 Vaccine in Healthcare Workers. Open Forum Infectious Diseases, 2022, 9, ofab656.	0.9	11
14	How Do Experts Treat Patients with BullousÂPemphigoid around the World? AnÂInternational Survey. JID Innovations, 2022, 2, 100129.	2.4	2
15	Severe cutaneous adverse reactions associated with systemic ivermectin: A pharmacovigilance analysis. Journal of Dermatology, 2022, 49, 769-774.	1.2	7
16	Coexistence of pachyonychia congenita and hidradenitis suppurativa: more than a coincidence. British Journal of Dermatology, 2022, 187, 392-400.	1.5	7
17	Heterozygous variants in the integrin subunit beta 4 gene (ITGB4) cause autosomal dominant nail dystrophy. British Journal of Dermatology, 2022, 187, 826-828.	1.5	1
18	A unique skin phenotype resulting from a large heterozygous deletion spanning six keratin genes. British Journal of Dermatology, 2022, 187, 773-777.	1.5	2

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19	Comorbidities in patients with palmoplantar plaque psoriasis. Journal of the American Academy of Dermatology, 2021, 84, 639-643.	1.2	8
20	Palmoplantar keratoderma caused by a missense variant in CTSB encoding cathepsin B. Clinical and Experimental Dermatology, 2021, 46, 103-108.	1.3	5
21	Laboratory monitoring during antifungal treatment of paediatric tinea capitis. Mycoses, 2021, 64, 157-161.	4.0	4
22	Molecular epidemiology of pachyonychia congenita in the Israeli population. Clinical and Experimental Dermatology, 2021, 46, 663-668.	1.3	1
23	ST18 affects cellâ€“cell adhesion in pemphigus vulgaris in a tumour necrosis factorâ€Î±â€dependent fashion*. British Journal of Dermatology, 2021, 184, 1153-1160.	1.5	7
24	The superiority of 72Âh leukocyte descent over CRP for mortality prediction in patients with sepsis. Clinica Chimica Acta, 2021, 514, 34-39.	1.1	4
25	Pulse-Dye Laser Followed by Betamethasone-Calcipotriol and Fractional Ablative CO2-Laser-Assisted Delivery for Nail Psoriasis. Dermatologic Surgery, 2021, 47, e111-e116.	0.8	11
26	Epidermolysis bullosa simplex due to biâ€allelic DST mutations: Case series and review of the literature. Pediatric Dermatology, 2021, 38, 436-441.	0.9	9
27	Atypical presentation of laryngoâ€onychoâ€cutaneous syndrome resulting from novel mutations in LAMA3A. Clinical and Experimental Dermatology, 2021, 46, 990-992.	1.3	0
28	Epidermolytic epidermal nevus caused by a somatic mutation in KRT2. Pediatric Dermatology, 2021, 38, 538-540.	0.9	1
29	Identification of clinically useful predictive genetic variants in pachyonychia congenita. Clinical and Experimental Dermatology, 2021, 46, 867-873.	1.3	5
30	Molecular epidemiology of nonâ€syndromic autosomal recessive congenital ichthyosis in a Middleâ€Eastern population. Experimental Dermatology, 2021, 30, 1290-1297.	2.9	10
31	Moisture Response Films Versus the Starch Iodine Test for the Detection of Palmar Hyperhidrosis. Dermatologic Surgery, 2021, Publish Ahead of Print, 668-671.	0.8	0
32	Laryngeal mucous membrane pemphigoid serves as a prognostic factor for poor response to treatment with rituximab. Clinical and Experimental Dermatology, 2021, 46, 915-919.	1.3	3
33	Diffuse Facial Hyperpigmentation as a Presenting Sign of Lupus Erythematosus: Three Cases and Review of the Literature. Case Reports in Dermatology, 2021, 13, 263-270.	0.8	4
34	Association Between Vaccination With BNT162b2 and Incidence of Symptomatic and Asymptomatic SARS-CoV-2 Infections Among Health Care Workers. JAMA - Journal of the American Medical Association, 2021, 325, 2457.	7.4	190
35	Primary Cutaneous B-Cell Lymphomas in Children and Adolescents: A SEER Population-Based Study. Clinical Lymphoma, Myeloma and Leukemia, 2021, 21, e1000-e1005.	0.4	6
36	Morphological features of benign pigmented ear lesions: a crossâ€sectional study. International Journal of Dermatology, 2021, , .	1.0	0

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37	Evidence for cutaneous dysbiosis in dystrophic epidermolysis bullosa. Clinical and Experimental Dermatology, 2021, 46, 1223-1229.	1.3	10
38	Ancestral patterns of recessive dystrophic epidermolysis bullosa mutations in Hispanic populations suggest sephardic ancestry. American Journal of Medical Genetics, Part A, 2021, 185, 3390-3400.	1.2	1
39	The pathogenesis of melasma and implications for treatment. Journal of Cosmetic Dermatology, 2021, 20, 3432-3445.	1.6	25
40	Role of Patch Testing in Chronic Spontaneous Urticaria. Journal of Asthma and Allergy, 2021, Volume 14, 1075-1079.	3.4	1
41	A splitâ€face clinical trial of conventional redâ€light photodynamic therapy versus daylight photodynamic therapy for acne vulgaris. Journal of Cosmetic Dermatology, 2021, 20, 3924-3930.	1.6	6
42	Immunogenicity of a BNT162b2 vaccine booster in health-care workers. Lancet Microbe, The, 2021, 2, e650.	7.3	20
43	Patch testing versus interferonâ€gamma release assay in evaluation of drug eruptions. Fundamental and Clinical Pharmacology, 2021, , .	1.9	4
44	Effect of a nationwide booster vaccine rollout in Israel on SARS-CoV-2 infection and severe illness in young adults. Travel Medicine and Infectious Disease, 2021, 44, 102195.	3.0	5
45	Syphilis Manifesting with Unilateral Hearing Loss and Tinnitus.. Indian Journal of Dermatology, 2021, 66, 575.	0.3	0
46	A treatment protocol for botulinum toxin injections in the treatment of pachyonychia congenitaâ€associated keratoderma. British Journal of Dermatology, 2020, 182, 671-677.	1.5	19
47	The Role of Desmoglein 1 in Gap Junction Turnover Revealed through the Study of SAMÂSyndrome. Journal of Investigative Dermatology, 2020, 140, 556-567.e9.	0.7	17
48	The combined effect of tranilast 8% liposomal gel on the final cosmesis of acne scarring in patients concomitantly treated by isotretinoin: prospective, doubleâ€blind, splitâ€face study. Clinical and Experimental Dermatology, 2020, 45, 41-47.	1.3	3
49	Coagulation Factor XIII-A Subunit Missense Mutation in the Pathobiology of Autosomal Dominant Multiple Dermatofibromas. Journal of Investigative Dermatology, 2020, 140, 624-635.e7.	0.7	12
50	Laser pretreatment for the attenuation of planned surgical scars: A randomized self-controlled hemi-scar pilot study. Journal of Plastic, Reconstructive and Aesthetic Surgery, 2020, 73, 893-898.	1.0	1
51	Treatment of epidermolysis bullosa pruriginosaâ€associated pruritus with dupilumab. British Journal of Dermatology, 2020, 182, 1495-1497.	1.5	41
52	Revisiting pachyonychia congenita: a caseâ€cohort study of 815 patients. British Journal of Dermatology, 2020, 182, 738-746.	1.5	31
53	An enhanced transcutaneous delivery of botulinum toxin for the treatment of Haileyâ€“Hailey disease. Dermatologic Therapy, 2020, 33, e13184.	1.7	6
54	Symptomatic mucosal involvement in pachyonychia congenita: challenges in infants and young children. British Journal of Dermatology, 2020, 182, 708-713.	1.5	5

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55	Treatment of hereditary hypotrichosis simplex of the scalp with topical gentamicin. British Journal of Dermatology, 2020, 183, 114-120.	1.5	19
56	Consensus reclassification of inherited epidermolysis bullosa and other disorders with skin fragility. British Journal of Dermatology, 2020, 183, 614-627.	1.5	406
57	Multidisciplinary care of epidermolysis bullosa during the COVID-19 pandemicâ€"Consensus: Recommendations by an international panel of experts. Journal of the American Academy of Dermatology, 2020, 83, 1222-1224.	1.2	7
58	Proximity Ligation Assay for Detecting Proteinâ€Protein Interactions and Protein Modifications in Cells and Tissues in Situ. Current Protocols in Cell Biology, 2020, 89, e115.	2.3	41
59	Phenotypic suppression of acral peeling skin syndrome in a patient with autosomal recessive congenital ichthyosis. Experimental Dermatology, 2020, 29, 742-748.	2.9	2
60	Updated S2K guidelines on the management of pemphigus vulgaris and foliaceus initiated by the european academy of dermatology and venereology (EADV). Journal of the European Academy of Dermatology and Venereology, 2020, 34, 1900-1913.	2.4	159
61	Loss-of-function variants in C3ORF52 result in localized autosomal recessive hypotrichosis. Genetics in Medicine, 2020, 22, 1227-1234.	2.4	12
62	Management Patterns of Delayed Inflammatory Reactions to Hyaluronic Acid Dermal Fillers: An Online Survey in Israel . Clinical, Cosmetic and Investigational Dermatology, 2020, Volume 13, 345-349.	1.8	9
63	Delayed Inflammatory Reactions to Hyaluronic Acid Fillers: A Literature Review and Proposed Treatment Algorithm . Clinical, Cosmetic and Investigational Dermatology, 2020, Volume 13, 371-378.	1.8	48
64	Intense focused ultrasound for neck and lower face skin tightening a prospective study. Journal of Cosmetic Dermatology, 2020, 19, 850-854.	1.6	9
65	Mucous membrane pemphigoidâ€"otorhinolaryngological manifestations: a retrospective cohort study. European Archives of Oto-Rhino-Laryngology, 2020, 277, 939-945.	1.6	9
66	Bullous pemphigoid distributed above the injury level in a paraplegic patient. Clinical and Experimental Dermatology, 2020, 45, 531-533.	1.3	1
67	Loss-of-Function Variants in SERPINA12 Underlie Autosomal Recessive Palmoplantar Keratoderma. Journal of Investigative Dermatology, 2020, 140, 2178-2187.	0.7	14
68	Identification of a founder mutation in <scp>KRT</scp> 14 associated with Naegeliâ€"Franceschettiâ€"Jadassohn syndrome. British Journal of Dermatology, 2020, 183, 756-757.	1.5	2
69	Transient Pruritic Erythema as a Forme Fruste of Solar Urticaria. Israel Medical Association Journal, 2020, 22, 227-231.	0.1	0
70	Griseofulvin vs terbinafine for paediatric tinea capitis: When and for how long. Mycoses, 2019, 62, 949-953.	4.0	14
71	PLACK syndrome shows remarkable phenotypic homogeneity. Clinical and Experimental Dermatology, 2019, 44, 580-583.	1.3	8
72	ST18 Enhances PV-IgG-Induced Loss of Keratinocyte Cohesion in Parallel to Increased ERK Activation. Frontiers in Immunology, 2019, 10, 770.	4.8	20

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73	Early intervention with pulse dye and CO2 ablative fractional lasers to improve cutaneous scarring post-lumpectomy: a randomized controlled trial on the impact of intervention on final cosmesis. Lasers in Medical Science, 2019, 34, 1881-1887.	2.1	11
74	Efficacy of a combination of diluted calcium hydroxylapatiteâ€based filler and an energyâ€based device for the treatment of facial atrophic acne scars. Clinical and Experimental Dermatology, 2019, 44, e171-e176.	1.3	9
75	Mutations in Recessive Congenital Ichthyoses Illuminate the Origin and Functions of the CorneocyteÂLipid Envelope. Journal of Investigative Dermatology, 2019, 139, 760-768.	0.7	41
76	Nested caseâ€“control study investigating the diagnostic role of tissue eosinophilia in adverse cutaneous drug reactions. Journal of the European Academy of Dermatology and Venereology, 2019, 33, 1152-1157.	2.4	4
77	Glutathione Sâ€transferase polymorphisms in patients with photosensitive and nonâ€photosensitive drug eruptions. Photodermatology Photoimmunology and Photomedicine, 2019, 35, 214-220.	1.5	4
78	Variant PADI3 in Central Centrifugal Cicatricial Alopecia. New England Journal of Medicine, 2019, 380, 833-841.	27.0	102
79	Effectiveness of topical propranolol 4% gel in the treatment of pyogenic granuloma in children. Journal of Dermatology, 2019, 46, 245-248.	1.2	11
80	Demodicidosis of the nipple. Lancet Infectious Diseases, The, 2019, 19, 112.	9.1	1
81	Loss-of-function mutations in caspase recruitment domain-containing protein 14 (CARD14) are associated with a severe variant of atopic dermatitis. Journal of Allergy and Clinical Immunology, 2019, 143, 173-181.e10.	2.9	60
82	Grover disease and bullous pemphigoid: a clinicopathological study of six cases. Clinical and Experimental Dermatology, 2019, 44, 524-527.	1.3	7
83	Anterior Scleritis Associated with Pemphigus Vulgaris. Ocular Immunology and Inflammation, 2019, 27, 497-498.	1.8	6
84	Comparative Study of Frozen and Paraffin-Embedded Sections: Evaluation of Inflammatory Dermatoses. Israel Medical Association Journal, 2019, 21, 82-84.	0.1	0
85	Successful treatment of Schambergâ€™s disease with fractional non-ablative 1540 nm erbium:glass laser. Journal of Cosmetic and Laser Therapy, 2018, 20, 265-268.	0.9	7
86	Topical pimecrolimus for paediatric cutaneous mastocytosis. Clinical and Experimental Dermatology, 2018, 43, 559-565.	1.3	17
87	Punctate palmoplantar keratoderma: an unusual mutation causing an unusual phenotype. British Journal of Dermatology, 2018, 178, 1455-1457.	1.5	5
88	NEK3-mediated SNAP29 phosphorylation modulates its membrane association and SNARE fusion dependent processes. Biochemical and Biophysical Research Communications, 2018, 497, 605-611.	2.1	7
89	Fractional ablative carbon dioxide laser followed by topical sodium stibogluconate application: A treatment option for pediatric cutaneous leishmaniasis. Pediatric Dermatology, 2018, 35, 366-369.	0.9	9
90	SAM syndrome is characterized by extensive phenotypic heterogeneity. Experimental Dermatology, 2018, 27, 787-790.	2.9	22

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91	Chronic pain in pachyonychia congenita: evidence for neuropathic origin. British Journal of Dermatology, 2018, 179, 154-162.	1.5	23
92	Striate palmoplantar keratoderma resulting from a missense mutation in DSG1. British Journal of Dermatology, 2018, 179, 755-757.	1.5	8
93	Novel POFUT 1 mutation associated with hidradenitis suppurativaâ€“Dowlingâ€“Degos disease firm up a role for Notch signalling in the pathogenesis of this disorder: reply from the authors. British Journal of Dermatology, 2018, 178, 986-986.	1.5	3
94	Recessive epidermolytic ichthyosis results from loss of keratin 10 expression, regardless of the mutation location. Clinical and Experimental Dermatology, 2018, 43, 187-190.	1.3	10
95	Immuneâ€regulatory genes as possible modifiers of familial pityriasis rubra pilaris â€“ lessons from a family with <scp>PRP</scp> and psoriasis. Journal of the European Academy of Dermatology and Venereology, 2018, 32, e389-e392.	2.4	6
96	Mechanisms Causing Loss of Keratinocyte Cohesion in Pemphigus. Journal of Investigative Dermatology, 2018, 138, 32-37.	0.7	113
97	A phenotype combining hidradenitis suppurativa with Dowling-Degos disease caused by a founder mutation in PSENEN. British Journal of Dermatology, 2018, 178, 502-508.	1.5	48
98	The Genetics of Pemphigus Vulgaris. Frontiers in Medicine, 2018, 5, 226.	2.6	60
99	Novel Stereoscopic Optical System for Objectively Measuring Above-Surface Scar Volumeâ€”First-Time Quantification of Responses to Various Treatment Modalities. Dermatologic Surgery, 2018, 44, 848-854.	0.8	10
100	Filaggrin 2 Deficiency Results in Abnormal Cell-Cell Adhesion in the Cornified Cell Layers and Causes Peeling Skin Syndrome Type A. Journal of Investigative Dermatology, 2018, 138, 1736-1743.	0.7	37
101	Identification of a recurrent mutation in ATP2C1 demonstrates that papular acantholytic dyskeratosis and Hailey-Hailey disease are allelic disorders. British Journal of Dermatology, 2018, 179, 1001-1002.	1.5	11
102	Bullous pemphigoid and diabetes mellitus: Are we missing the larger picture?. Journal of the American Academy of Dermatology, 2018, 79, e27.	1.2	6
103	Assessment of the effectiveness of topical propranolol 4% gel for infantile hemangiomas. International Journal of Dermatology, 2017, 56, 148-153.	1.0	18
104	Occupational mycosis fungoides â€“ a case series. International Journal of Dermatology, 2017, 56, 733-737.	1.0	7
105	The Molecular Revolution in Cutaneous Biology: EraÂof Next-Generation Sequencing. Journal of Investigative Dermatology, 2017, 137, e79-e82.	0.7	16
106	A distinct cutaneous microbiota profile in autoimmune bullous disease patients. Experimental Dermatology, 2017, 26, 1221-1227.	2.9	28
107	Image Gallery: Massive localized lymphoedema. British Journal of Dermatology, 2017, 176, e95-e95.	1.5	1
108	IgA pemphigus: lumping or splitting?. British Journal of Dermatology, 2017, 177, 581-582.	1.5	2

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109	Meeting Report of the Pathogenesis of Pemphigus and Pemphigoid Meeting in Munich, September 2016. Journal of Investigative Dermatology, 2017, 137, 1199-1203.	0.7	34
110	Report of the 13th Annual International Pachyonychia Congenita Consortium Symposium. British Journal of Dermatology, 2017, 176, 1144-1147.	1.5	6
111	Predicting neurofibromatosis type 1 riskÂamong children with isolated cafÃ©-au-laitÂmacules. Journal of the American Academy of Dermatology, 2017, 76, 1077-1083.e3.	1.2	28
112	ARCI7 Revisited and Repositioned. Journal of Investigative Dermatology, 2017, 137, 970-972.	0.7	6
113	Paraneoplastic pityriasis rubra pilaris: case report and literature review. Clinical and Experimental Dermatology, 2017, 42, 54-57.	1.3	14
114	Failure of initial disease control in bullous pemphigoid: a retrospective study of hospitalized patients in a single tertiary center. International Journal of Dermatology, 2017, 56, 1010-1016.	1.0	7
115	Rituximab and shortâ€course prednisone as the new gold standard for newâ€onset pemphigus vulgaris and pemphigus foliaceus. British Journal of Dermatology, 2017, 177, 1143-1144.	1.5	14
116	The Effect of Tranilast 8% Liposomal Gel Versus Placebo on Post-Cesarean Surgical Scars: A Prospective Double-Blind Split-Scar Study. Dermatologic Surgery, 2017, 43, 1157-1163.	0.8	6
117	Calpain 12 Function Revealed through the Study of an Atypical Case of Autosomal Recessive Congenital Ichthyosis. Journal of Investigative Dermatology, 2017, 137, 385-393.	0.7	19
118	<scp>SVEP</scp>1 plays a crucial role in epidermal differentiation. Experimental Dermatology, 2017, 26, 423-430.	2.9	17
119	Giant pyogenic granuloma of theÂfinger in an HIVâ€positive patient. Journal of the European Academy of Dermatology and Venereology, 2017, 31, e512-e513.	2.4	0
120	NB-UVB phototherapy for generalized granuloma annulare. Dermatologic Therapy, 2016, 29, 152-154.	1.7	11
121	Tinea capitis outbreak among paediatric refugee population, an evolving healthcare challenge. Mycoses, 2016, 59, 553-557.	4.0	31
122	Isotretinoin treatment of autosomal recessive congenital ichthyosis complicated by coexisting dysferlinopathy. Clinical and Experimental Dermatology, 2016, 41, 390-393.	1.3	5
123	Segmental basal cell naevus syndrome caused by an activating mutation in smoothened. British Journal of Dermatology, 2016, 175, 178-181.	1.5	33
124	A refractory, cutaneous, subepidermal bullous disease. Clinical and Experimental Dermatology, 2016, 41, 573-575.	1.3	6
125	Understanding unspecific complaints through genetics. Nature Genetics, 2016, 48, 1450-1451.	21.4	0
126	Pemphigoid: diversity in evolution. British Journal of Dermatology, 2016, 175, 676-677.	1.5	2

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127	Loss-of-Function Mutations in SERPINB8 Linked to Exfoliative Ichthyosis with Impaired Mechanical Stability of Intercellular Adhesions. American Journal of Human Genetics, 2016, 99, 430-436.	6.2	27
128	Nonâ€keratinocyte <scp>SNAP</scp>29 influences epidermal differentiation and hair follicle formation in mice. Experimental Dermatology, 2016, 25, 647-649.	2.9	8
129	Monopathogenic vs multipathogenic explanations of pemphigus pathophysiology. Experimental Dermatology, 2016, 25, 839-846.	2.9	63
130	Mutations in Three Genes Encoding Proteins Involved in Hair Shaft Formation Cause Uncombable Hair Syndrome. American Journal of Human Genetics, 2016, 99, 1292-1304.	6.2	127
131	Happle-Tinschert syndrome can be caused by a mosaicSMOmutation and is suggested to be a variant of Curry-Jones syndrome: reply from the authors. British Journal of Dermatology, 2016, 175, 1109-1109.	1.5	4
132	A novel homozygous deletion inEXPH5causes a skin fragility phenotype. Clinical and Experimental Dermatology, 2016, 41, 915-918.	1.3	7
133	Stabilizing mutations of KLHL24 ubiquitin ligase cause loss of keratin 14 and human skin fragility. Nature Genetics, 2016, 48, 1508-1516.	21.4	101
134	Somatic Mosaicism for a â€œLethalâ€•<scp>GJB</scp>2 Mutation Results in a Patterned Form of Spiny Hyperkeratosis without Eccrine Involvement. Pediatric Dermatology, 2016, 33, 322-326.	0.9	5
135	Papillonâ€“LefÃ¨vre syndrome: report of six patients and identification of a novel mutation. International Journal of Dermatology, 2016, 55, 898-902.	1.0	11
136	Establishment of Two Mouse Models for CEDNIK Syndrome Reveals the Pivotal Role of SNAP29 in Epidermal Differentiation. Journal of Investigative Dermatology, 2016, 136, 672-679.	0.7	31
137	Childhood Pemphigus Foliaceus with Exclusive Immunoglobulin G Autoantibodies to Desmocollins. Pediatric Dermatology, 2016, 33, e10-3.	0.9	9
138	Identification of a Functional Risk Variant for Pemphigus Vulgaris in the ST18 Gene. PLoS Genetics, 2016, 12, e1006008.	3.5	53
139	Mutations in TSPEAR, Encoding a Regulator of Notch Signaling, Affect Tooth and Hair Follicle Morphogenesis. PLoS Genetics, 2016, 12, e1006369.	3.5	32
140	BJDin translation. British Journal of Dermatology, 2015, 173, 1349-1350.	1.5	0
141	Paraneoplastic pemphigus: an entity still in search of an identity?. British Journal of Dermatology, 2015, 173, 1363-1364.	1.5	3
142	Novel <scp>TGM</scp>5 mutations in acral peeling skin syndrome. Experimental Dermatology, 2015, 24, 285-289.	2.9	11
143	Autosomalâ€dominant cutis laxa resulting from an intronic mutation in <scp>ELN</scp>. Experimental Dermatology, 2015, 24, 885-887.	2.9	4
144	Definitions and outcome measures for mucous membrane pemphigoid: Recommendations ofÂanÂinternational panel of experts. Journal of the American Academy of Dermatology, 2015, 72, 168-174.	1.2	133

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145	Extensive lentigo simplex, linear epidermolytic naevus and epidermolytic naevus comedonicus caused by a somatic mutation inKRT10. British Journal of Dermatology, 2015, 173, 293-296.	1.5	9
146	Inherited desmosomal disorders. Cell and Tissue Research, 2015, 360, 457-475.	2.9	29
147	Severe dermatitis, multiple allergies, and metabolic wasting syndrome caused by a novel mutation in the N-terminal plakin domain of desmoplakin. Journal of Allergy and Clinical Immunology, 2015, 136, 1268-1276.	2.9	103
148	Middle Cerebral Artery Stenosis in Patients with Acute Ischemic Stroke and TIA in Israel. American Journal of Neuroradiology, 2015, 36, 46-49.	2.4	7
149	Pyoderma gangrenosum, acne and ulcerative colitis in a patient with a novel mutation in thePSTPIP1gene. Clinical and Experimental Dermatology, 2015, 40, 367-372.	1.3	53
150	Analysis of CARD14 Polymorphisms in Pityriasis Rubra Pilaris: Activation of NF-ÎºB. Journal of Investigative Dermatology, 2015, 135, 1905-1908.	0.7	24
151	A case for diagnosis. Clinical and Experimental Dermatology, 2015, 40, 697-699.	1.3	1
152	<scp>RBM</scp>28, a protein deficient in <scp>ANE</scp> syndrome, regulates hair follicle growth via miRâ€203 and p63. Experimental Dermatology, 2015, 24, 618-622.	2.9	17
153	Mutations inSMARCAD1cause autosomal dominant adermatoglyphia and perturb the expression of epidermal differentiation-associated genes. British Journal of Dermatology, 2014, 171, 1521-1524.	1.5	16
154	A Mutation in TP63 Causing a Mild Ectodermal Dysplasia Phenotype. Journal of Investigative Dermatology, 2014, 134, 2277-2280.	0.7	5
155	Increased epidermal expression and absence of mutations inCARD14in a series of patients with sporadic pityriasis rubra pilaris. British Journal of Dermatology, 2014, 170, 1196-1198.	1.5	21
156	Disease classification using clinical and molecular features. Journal of Dermatology, 2014, 41, 949-949.	1.2	0
157	The molecular genetic analysis of the expanding pachyonychia congenita case collection. British Journal of Dermatology, 2014, 171, 343-355.	1.5	54
158	What's in a disease name?. British Journal of Dermatology, 2014, 170, 1005-1007.	1.5	3
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160	Olmsted Syndrome Caused by a Homozygous Recessive Mutation in TRPV3. Journal of Investigative Dermatology, 2014, 134, 1752-1754.	0.7	44
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