Eli Sprecher

List of Publications by Year in Descending Order

220 papers	5,873 citations	40 h-index	70 g-index
241 ext. papers	7,296 ext. citations	4.5 avg, IF	5.66 L-index

#	Paper	IF	Citations
220	Neonatal inflammatory skin and bowel disease type 1 caused by a complex genetic defect and responsive to combined anti-TNF-alpha and IL-12/23 blockade.. *British Journal of Dermatology*, **2022** ,	4	0
219	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**,	27.4	14
218	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	4.9	0
217	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	1.1	
216	Short-Term Safety of Booster Immunization With BNT162b2 mRNA COVID-19 Vaccine in Healthcare Workers.. *Open Forum Infectious Diseases*, **2022**, 9, ofab656	1	0
215	How do experts treat patients with bullous pemphigoid around the world? An international survey.. *JID Innovations*, **2022**, 100129		
214	Severe cutaneous adverse reactions associated with systemic ivermectin: A pharmacovigilance analysis.. *Journal of Dermatology*, **2022**,	1.6	1
213	Translational implications of Th17-skewed inflammation due to genetic deficiency of a cadherin stress sensor.. *Journal of Clinical Investigation*, **2021**,	15.9	4
212	Immunogenicity of a BNT162b2 vaccine booster in health-care workers. *Lancet Microbe, The*, **2021**, 2, e650	22.2	9
211	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	8.4	3
210	Molecular epidemiology of non-syndromic autosomal recessive congenital ichthyosis in a Middle-Eastern population. *Experimental Dermatology*, **2021**, 30, 1290-1297	4	3
209	Moisture Response Films Versus the Starch Iodine Test for the Detection of Palmar Hyperhidrosis. *Dermatologic Surgery*, **2021**, 47, 668-671	1.7	
208	Laryngeal mucous membrane pemphigoid serves as a prognostic factor for poor response to treatment with rituximab. *Clinical and Experimental Dermatology*, **2021**, 46, 915-919	1.8	1
207	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	1.1	0
206	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, 2457-2465	27.4	111
205	Primary Cutaneous B-Cell Lymphomas in Children and Adolescents: A SEER Population-Based Study. *Clinical Lymphoma, Myeloma and Leukemia*, **2021**, 21, e1000-e1005	2	0
204	Comorbidities in patients with palmoplantar plaque psoriasis. *Journal of the American Academy of Dermatology*, **2021**, 84, 639-643	4.5	4

203	Palmoplantar keratoderma caused by a missense variant in CTSB encoding cathepsin B. *Clinical and Experimental Dermatology*, **2021**, 46, 103-108	1.8	1
202	Laboratory monitoring during antifungal treatment of paediatric tinea capitis. *Mycoses*, **2021**, 64, 157-161	3.2	1
201	Molecular epidemiology of pachyonychia congenita in the Israeli population. *Clinical and Experimental Dermatology*, **2021**, 46, 663-668	1.8	0
200	ST18 affects cell-cell adhesion in pemphigus vulgaris in a tumour necrosis factor-□dependent fashion. *British Journal of Dermatology*, **2021**, 184, 1153-1160	4	6
199	The superiority of 72 h leukocyte descent over CRP for mortality prediction in patients with sepsis. *Clinica Chimica Acta*, **2021**, 514, 34-39	6.2	1
198	Pulse-Dye Laser Followed by Betamethasone-Calcipotriol and Fractional Ablative CO2-Laser-Assisted Delivery for Nail Psoriasis. *Dermatologic Surgery*, **2021**, 47, e111-e116	1.7	5
197	Epidermolysis bullosa simplex due to bi-allelic DST mutations: Case series and review of the literature. *Pediatric Dermatology*, **2021**, 38, 436-441	1.9	2
196	Atypical presentation of laryngo-onycho-cutaneous syndrome resulting from novel mutations in LAMA3A. *Clinical and Experimental Dermatology*, **2021**, 46, 990-992	1.8	
195	Laryngeal Pemphigoid Evolution and Response to Treatment. *Journal of Voice*, **2021**,	1.9	1
194	Epidermolytic epidermal nevus caused by a somatic mutation in KRT2. *Pediatric Dermatology*, **2021**, 38, 538-540	1.9	1
193	Identification of clinically useful predictive genetic variants in pachyonychia congenita. *Clinical and Experimental Dermatology*, **2021**, 46, 867-873	1.8	0
192	Evidence for cutaneous dysbiosis in dystrophic epidermolysis bullosa. *Clinical and Experimental Dermatology*, **2021**, 46, 1223-1229	1.8	2
191	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	2.5	
190	The pathogenesis of melasma and implications for treatment. *Journal of Cosmetic Dermatology*, **2021**, 20, 3432-3445	2.5	6
189	Role of Patch Testing in Chronic Spontaneous Urticaria. *Journal of Asthma and Allergy*, **2021**, 14, 1075-1079	3.1	
188	A split-face clinical trial of conventional red-light photodynamic therapy versus daylight photodynamic therapy for acne vulgaris. *Journal of Cosmetic Dermatology*, **2021**, 20, 3924-3930	2.5	0
187	Clinical efficacy of fecal microbial transplantation treatment in adults with moderate-to-severe atopic dermatitis.. *Immunity, Inflammation and Disease*, **2021**,	2.4	4
186	Syphilis Manifesting with Unilateral Hearing Loss and Tinnitus.. *Indian Journal of Dermatology*, **2021**, 66, 575	0.9	

185	Loss-of-function variants in C3ORF52 result in localized autosomal recessive hypotrichosis. *Genetics in Medicine*, **2020**, 22, 1227-1234	8.1	6
184	Management Patterns of Delayed Inflammatory Reactions to Hyaluronic Acid Dermal Fillers: An Online Survey in Israel. *Clinical, Cosmetic and Investigational Dermatology*, **2020**, 13, 345-349	2.9	3
183	Delayed Inflammatory Reactions to Hyaluronic Acid Fillers: A Literature Review and Proposed Treatment Algorithm. *Clinical, Cosmetic and Investigational Dermatology*, **2020**, 13, 371-378	2.9	19
182	Intense focused ultrasound for neck and lower face skin tightening a prospective study. *Journal of Cosmetic Dermatology*, **2020**, 19, 850-854	2.5	4
181	Mucous membrane pemphigoid-otorhinolaryngological manifestations: a retrospective cohort study. *European Archives of Oto-Rhino-Laryngology*, **2020**, 277, 939-945	3.5	6
180	Bullous pemphigoid distributed above the injury level in a paraplegic patient. *Clinical and Experimental Dermatology*, **2020**, 45, 531-533	1.8	
179	Loss-of-Function Variants in SERPINA12 Underlie Autosomal Recessive Palmoplantar Keratoderma. *Journal of Investigative Dermatology*, **2020**, 140, 2178-2187	4.3	4
178	Coagulation Factor XIII-A Subunit Missense Mutation in the Pathobiology of Autosomal Dominant Multiple Dermatofibromas. *Journal of Investigative Dermatology*, **2020**, 140, 624-635.e7	4.3	4
177	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	1.7	1
176	Treatment of epidermolysis bullosa pruriginosa-associated pruritus with dupilumab. *British Journal of Dermatology*, **2020**, 182, 1495-1497	4	25
175	Revisiting pachyonychia congenita: a case-cohort study of 815 patients. *British Journal of Dermatology*, **2020**, 182, 738-746	4	16
174	An enhanced transcutaneous delivery of botulinum toxin for the treatment of Hailey-Hailey disease. *Dermatologic Therapy*, **2020**, 33, e13184	2.2	4
173	Symptomatic mucosal involvement in pachyonychia congenita: challenges in infants and young children. *British Journal of Dermatology*, **2020**, 182, 708-713	4	2
172	Treatment of hereditary hypotrichosis simplex of the scalp with topical gentamicin. *British Journal of Dermatology*, **2020**, 183, 114-120	4	10
171	Consensus reclassification of inherited epidermolysis bullosa and other disorders with skin fragility. *British Journal of Dermatology*, **2020**, 183, 614-627	4	179
170	Multidisciplinary care of epidermolysis bullosa during the COVID-19 pandemic-Consensus: Recommendations by an international panel of experts. *Journal of the American Academy of Dermatology*, **2020**, 83, 1222-1224	4.5	5
169	Proximity Ligation Assay for Detecting Protein-Protein Interactions and Protein Modifications in Cells and Tissues in Situ. *Current Protocols in Cell Biology*, **2020**, 89, e115	2.3	8
168	Phenotypic suppression of acral peeling skin syndrome in a patient with autosomal recessive congenital ichthyosis. *Experimental Dermatology*, **2020**, 29, 742-748	4	0

167	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	4.6	60
166	A treatment protocol for botulinum toxin injections in the treatment of pachyonychia congenita-associated keratoderma. *British Journal of Dermatology*, **2020**, 182, 671-677	4	11
165	The Role of Desmoglein 1 in Gap Junction Turnover Revealed through the Study of SAM Syndrome. *Journal of Investigative Dermatology*, **2020**, 140, 556-567.e9	4.3	8
164	The combined effect of tranilast 8% liposomal gel on the final cosmesis of acne scarring in patients concomitantly treated by isotretinoin: prospective, double-blind, split-face study. *Clinical and Experimental Dermatology*, **2020**, 45, 41-47	1.8	2
163	Identification of a founder mutation in KRT14 associated with Naegeli-Franceschetti-Jadassohn syndrome. *British Journal of Dermatology*, **2020**, 183, 756-757	4	1
162	Transient Pruritic Erythema as a Forme Fruste of Solar Urticaria. *Israel Medical Association Journal*, **2020**, 22, 227-231	0.9	
161	PLACK syndrome shows remarkable phenotypic homogeneity. *Clinical and Experimental Dermatology*, **2019**, 44, 580-583	1.8	4
160	ST18 Enhances PV-IgG-Induced Loss of Keratinocyte Cohesion in Parallel to Increased ERK Activation. *Frontiers in Immunology*, **2019**, 10, 770	8.4	11
159	Early intervention with pulse dye and CO 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	3.1	8
158	Efficacy of a combination of diluted calcium hydroxylapatite-based filler and an energy-based device for the treatment of facial atrophic acne scars. *Clinical and Experimental Dermatology*, **2019**, 44, e171-e176	1.8	6
157	Mutations in Recessive Congenital Ichthyoses Illuminate the Origin and Functions of the Corneocyte Lipid Envelope. *Journal of Investigative Dermatology*, **2019**, 139, 760-768	4.3	26
156	Nested case-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	4.6	3
155	Glutathione S-transferase polymorphisms in patients with photosensitive and non-photosensitive drug eruptions. *Photodermatology Photoimmunology and Photomedicine*, **2019**, 35, 214-220	2.4	4
154	Griseofulvin vs terbinafine for paediatric tinea capitis: When and for how long. *Mycoses*, **2019**, 62, 949-953	2	8
153	Variant in Central Centrifugal Cicatricial Alopecia. *New England Journal of Medicine*, **2019**, 380, 833-841	59.2	49
152	Dyschromatosis **2019**, 1499-1514		
151	Pityriasis Rubra Pilaris **2019**, 377-389		
150	Effectiveness of topical propranolol 4% gel in the treatment of pyogenic granuloma in children. *Journal of Dermatology*, **2019**, 46, 245-248	1.6	6

149	Demodicidosis of the nipple. *Lancet Infectious Diseases, The*, **2019**, 19, 112	25.5	0
148	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	11.5	33
147	Grover disease and bullous pemphigoid: a clinicopathological study of six cases. *Clinical and Experimental Dermatology*, **2019**, 44, 524-527	1.8	4
146	Anterior Scleritis Associated with Pemphigus Vulgaris. *Ocular Immunology and Inflammation*, **2019**, 27, 497-498	2.8	4
145	Comparative Study of Frozen and Paraffin-Embedded Sections: Evaluation of Inflammatory Dermatoses. *Israel Medical Association Journal*, **2019**, 21, 82-84	0.9	
144	Successful treatment of Schamberg's disease with fractional non-ablative 1540 nm erbium:glass laser. *Journal of Cosmetic and Laser Therapy*, **2018**, 20, 265-268	1.8	4
143	Topical pimecrolimus for paediatric cutaneous mastocytosis. *Clinical and Experimental Dermatology*, **2018**, 43, 559-565	1.8	11
142	Punctate palmoplantar keratoderma: an unusual mutation causing an unusual phenotype. *British Journal of Dermatology*, **2018**, 178, 1455-1457	4	4
141	NEK3-mediated SNAP29 phosphorylation modulates its membrane association and SNARE fusion dependent processes. *Biochemical and Biophysical Research Communications*, **2018**, 497, 605-611	3.4	5
140	Fractional ablative carbon dioxide laser followed by topical sodium stibogluconate application: A treatment option for pediatric cutaneous leishmaniasis. *Pediatric Dermatology*, **2018**, 35, 366-369	1.9	7
139	SAM syndrome is characterized by extensive phenotypic heterogeneity. *Experimental Dermatology*, **2018**, 27, 787-790	4	15
138	Chronic pain in pachyonychia congenita: evidence for neuropathic origin. *British Journal of Dermatology*, **2018**, 179, 154-162	4	16
137	Striate palmoplantar keratoderma resulting from a missense mutation in DSG1. *British Journal of Dermatology*, **2018**, 179, 755-757	4	7
136	Novel POFUT1 mutation associated with hidradenitis suppurativa-Dowling-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	4	2
135	Recessive epidermolytic ichthyosis results from loss of keratin 10 expression, regardless of the mutation location. *Clinical and Experimental Dermatology*, **2018**, 43, 187-190	1.8	7
134	Immune-regulatory genes as possible modifiers of familial pityriasis rubra pilaris - lessons from a family with PRP and psoriasis. *Journal of the European Academy of Dermatology and Venereology*, **2018**, 32, e389-e392	4.6	3
133	Mechanisms Causing Loss of Keratinocyte Cohesion in Pemphigus. *Journal of Investigative Dermatology*, **2018**, 138, 32-37	4.3	70
132	A phenotype combining hidradenitis suppurativa with Dowling-Degos disease caused by a founder mutation in PSENEN. *British Journal of Dermatology*, **2018**, 178, 502-508	4	35

131	Bullous pemphigoid and diabetes mellitus: Are we missing the larger picture?. *Journal of the American Academy of Dermatology*, **2018**, 79, e27	4.5	4
130	The Genetics of Pemphigus Vulgaris. *Frontiers in Medicine*, **2018**, 5, 226	4.9	36
129	Novel Stereoscopic Optical System for Objectively Measuring Above-Surface Scar Volume-First-Time Quantification of Responses to Various Treatment Modalities. *Dermatologic Surgery*, **2018**, 44, 848-854	1.7	5
128	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	4.3	23
127	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	4	7
126	Assessment of the effectiveness of topical propranolol 4% gel for infantile hemangiomas. *International Journal of Dermatology*, **2017**, 56, 148-153	1.7	15
125	Occupational mycosis fungoides - a case series. *International Journal of Dermatology*, **2017**, 56, 733-737	1.7	4
124	The Molecular Revolution in Cutaneous Biology: Era´of Next-Generation Sequencing. *Journal of Investigative Dermatology*, **2017**, 137, e79-e82	4.3	8
123	A distinct cutaneous microbiota profile in autoimmune bullous disease patients. *Experimental Dermatology*, **2017**, 26, 1221-1227	4	15
122	Image Gallery: Massive localized lymphoedema. *British Journal of Dermatology*, **2017**, 176, e95	4	1
121	IgA pemphigus: lumping or splitting?. *British Journal of Dermatology*, **2017**, 177, 581-582	4	1
120	Meeting Report of the Pathogenesis of Pemphigus and Pemphigoid Meeting in Munich, September 2016. *Journal of Investigative Dermatology*, **2017**, 137, 1199-1203	4.3	26
119	Report of the 13th Annual International Pachyonychia Congenita Consortium Symposium. *British Journal of Dermatology*, **2017**, 176, 1144-1147	4	4
118	Predicting neurofibromatosis type 1 risk´among children with isolated caf□au-lait´macules. *Journal of the American Academy of Dermatology*, **2017**, 76, 1077-1083.e3	4.5	20
117	ARCI7 Revisited and Repositioned. *Journal of Investigative Dermatology*, **2017**, 137, 970-972	4.3	3
116	Paraneoplastic pityriasis rubra pilaris: case report and literature review. *Clinical and Experimental Dermatology*, **2017**, 42, 54-57	1.8	12
115	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	1.7	6
114	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	1.7	5

113	Calpain 12 Function Revealed through the Study of an Atypical Case of Autosomal Recessive Congenital Ichthyosis. *Journal of Investigative Dermatology*, **2017**, 137, 385-393	4.3	14
112	SVEP1 plays a crucial role in epidermal differentiation. *Experimental Dermatology*, **2017**, 26, 423-430	4	6
111	Giant pyogenic granuloma of the finger in an HIV-positive patient. *Journal of the European Academy of Dermatology and Venereology*, **2017**, 31, e512-e513	4.6	
110	Monopathogenic vs multipathogenic explanations of pemphigus pathophysiology. *Experimental Dermatology*, **2016**, 25, 839-846	4	48
109	Mutations in Three Genes Encoding Proteins Involved in Hair Shaft Formation Cause Uncombable Hair Syndrome. *American Journal of Human Genetics*, **2016**, 99, 1292-1304	11	72
108	Happle-Tinschert syndrome can be caused by a mosaic SMO mutation and is suggested to be a variant of Curry-Jones syndrome: reply from the authors. *British Journal of Dermatology*, **2016**, 175, 1109	4	3
107	A novel homozygous deletion in EXPH5 causes a skin fragility phenotype. *Clinical and Experimental Dermatology*, **2016**, 41, 915-918	1.8	4
106	Stabilizing mutations of KLHL24 ubiquitin ligase cause loss of keratin 14 and human skin fragility. *Nature Genetics*, **2016**, 48, 1508-1516	36.3	73
105	Somatic Mosaicism for a "Lethal" GJB2 Mutation Results in a Patterned Form of Spiny Hyperkeratosis without Eccrine Involvement. *Pediatric Dermatology*, **2016**, 33, 322-6	1.9	2
104	Papillon-Lefvre syndrome: report of six patients and identification of a novel mutation. *International Journal of Dermatology*, **2016**, 55, 898-902	1.7	9
103	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	4.3	24
102	Childhood Pemphigus Foliaceus with Exclusive Immunoglobulin G Autoantibodies to Desmocollins. *Pediatric Dermatology*, **2016**, 33, e10-3	1.9	6
101	Identification of a Functional Risk Variant for Pemphigus Vulgaris in the ST18 Gene. *PLoS Genetics*, **2016**, 12, e1006008	6	38
100	Mutations in TSPEAR, Encoding a Regulator of Notch Signaling, Affect Tooth and Hair Follicle Morphogenesis. *PLoS Genetics*, **2016**, 12, e1006369	6	20
99	NB-UVB phototherapy for generalized granuloma annulare. *Dermatologic Therapy*, **2016**, 29, 152-4	2.2	8
98	Tinea capitis outbreak among paediatric refugee population, an evolving healthcare challenge. *Mycoses*, **2016**, 59, 553-7	5.2	26
97	Isotretinoin treatment of autosomal recessive congenital ichthyosis complicated by coexisting dysferlinopathy. *Clinical and Experimental Dermatology*, **2016**, 41, 390-3	1.8	2
96	Segmental basal cell naevus syndrome caused by an activating mutation in smoothened. *British Journal of Dermatology*, **2016**, 175, 178-81	4	22

95	A refractory, cutaneous, subepidermal bullous disease. *Clinical and Experimental Dermatology*, **2016**, 41, 573-5	1.8	5
94	Understanding unspecific complaints through genetics. *Nature Genetics*, **2016**, 48, 1450-1451	36.3	
93	Pemphigoid: diversity in evolution. *British Journal of Dermatology*, **2016**, 175, 676-7	4	1
92	Loss-of-Function Mutations in SERPINB8 Linked to Exfoliative Ichthyosis with Impaired Mechanical Stability of Intercellular Adhesions. *American Journal of Human Genetics*, **2016**, 99, 430-6	11	14
91	Non-keratinocyte SNAP29 influences epidermal differentiation and hair follicle formation in mice. *Experimental Dermatology*, **2016**, 25, 647-9	4	6
90	Inherited desmosomal disorders. *Cell and Tissue Research*, **2015**, 360, 457-75	4.2	25
89	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, 1268-76	11.5	82
88	Middle cerebral artery stenosis in patients with acute ischemic stroke and TIA in Israel. *American Journal of Neuroradiology*, **2015**, 36, 46-9	4.4	5
87	Pyoderma gangrenosum, acne and ulcerative colitis in a patient with a novel mutation in the PSTPIP1 gene. *Clinical and Experimental Dermatology*, **2015**, 40, 367-72	1.8	45
86	Analysis of CARD14 Polymorphisms in Pityriasis Rubra Pilaris: Activation of NF-Β. *Journal of Investigative Dermatology*, **2015**, 135, 1905-1908	4.3	20
85	A case for diagnosis. *Clinical and Experimental Dermatology*, **2015**, 40, 697-9	1.8	1
84	RBM28, a protein deficient in ANE syndrome, regulates hair follicle growth via miR-203 and p63. *Experimental Dermatology*, **2015**, 24, 618-22	4	13
83	Paraneoplastic pemphigus: an entity still in search of an identity?. *British Journal of Dermatology*, **2015**, 173, 1363-4	4	3
82	Novel TGM5 mutations in acral peeling skin syndrome. *Experimental Dermatology*, **2015**, 24, 285-9	4	10
81	Autosomal-dominant cutis laxa resulting from an intronic mutation in ELN. *Experimental Dermatology*, **2015**, 24, 885-7	4	2
80	Definitions and outcome measures for mucous membrane pemphigoid: recommendations of an international panel of experts. *Journal of the American Academy of Dermatology*, **2015**, 72, 168-74	4.5	93
79	Extensive lentigo simplex, linear epidermolytic naevus and epidermolytic naevus comedonicus caused by a somatic mutation in KRT10. *British Journal of Dermatology*, **2015**, 173, 293-6	4	7
78	The expanding spectrum of IgA pemphigus: a case report and review of the literature. *British Journal of Dermatology*, **2014**, 171, 650-6	4	25

77	Clinico-pathological manifestations of variant late infantile neuronal ceroid lipofuscinosis (vLINCL) caused by a novel mutation in MFSD8 gene. *European Journal of Medical Genetics*, **2014**, 57, 607-12	2.6	11
76	Peeling off the genetics of atopic dermatitis-like congenital disorders. *Journal of Allergy and Clinical Immunology*, **2014**, 134, 808-15	11.5	39
75	Angiomodulin is required for cardiogenesis of embryonic stem cells and is maintained by a feedback loop network of p63 and Activin-A. *Stem Cell Research*, **2014**, 12, 49-59	1.6	8
74	Autosomal dominant inheritance of central centrifugal cicatricial alopecia in black South Africans. *Journal of the American Academy of Dermatology*, **2014**, 70, 679-682.e1	4.5	41
73	Molecular analysis of a series of Israeli families with Com Netherton syndrome. *Dermatology*, **2014**, 228, 183-8	4.4	8
72	Mutations in SMARCAD1 cause autosomal dominant adermatoglyphia and perturb the expression of epidermal differentiation-associated genes. *British Journal of Dermatology*, **2014**, 171, 1521-4	4	12
71	A mutation in TP63 causing a mild ectodermal dysplasia phenotype. *Journal of Investigative Dermatology*, **2014**, 134, 2277-2280	4.3	2
70	Increased epidermal expression and absence of mutations in CARD14 in a series of patients with sporadic pityriasis rubra pilaris. *British Journal of Dermatology*, **2014**, 170, 1196-8	4	13
69	Disease classification using clinical and molecular features. *Journal of Dermatology*, **2014**, 41, 949	1.6	
68	The molecular genetic analysis of the expanding pachyonychia congenita case collection. *British Journal of Dermatology*, **2014**, 171, 343-55	4	48
67	Clinical response to ustekinumab in familial pityriasis rubra pilaris caused by a novel mutation in CARD14. *British Journal of Dermatology*, **2014**, 171, 420-2	4	48
66	Olmsted syndrome caused by a homozygous recessive mutation in TRPV3. *Journal of Investigative Dermatology*, **2014**, 134, 1752-1754	4.3	32
65	A novel splice-site mutation in the AAGAB gene segregates with hereditary punctate palmoplantar keratoderma and congenital dysplasia of the hip in a large family. *Clinical and Experimental Dermatology*, **2014**, 39, 182-6	1.8	9
64	Pachyonychia congenita cornered: report on the 11th Annual International Pachyonychia Congenita Consortium Meeting. *British Journal of Dermatology*, **2014**, 171, 974-7	4	9
63	Best treatment practices for pachyonychia congenita. *Journal of the European Academy of Dermatology and Venereology*, **2014**, 28, 279-85	4.6	30
62	Topobiology of human pigmentation: P-cadherin selectively stimulates hair follicle melanogenesis. *Journal of Investigative Dermatology*, **2013**, 133, 1591-600	4.3	22
61	Cole Disease Results from Mutations in ENPP1. *American Journal of Human Genetics*, **2013**, 93, 752-7	11	28
60	Semidominant inheritance in epidermolytic ichthyosis. *Journal of Investigative Dermatology*, **2013**, 133, 2626-2628	4.3	7

59	Non-syndromic autosomal recessive congenital ichthyosis in the Israeli population. *Clinical and Experimental Dermatology*, **2013**, 38, 911-6	1.8	24
58	Desmoglein 1 deficiency results in severe dermatitis, multiple allergies and metabolic wasting. *Nature Genetics*, **2013**, 45, 1244-1248	36.3	217
57	Comparative study of high-resolution multifrequency ultrasound of the plantar skin in patients with various types of hereditary palmoplantar keratoderma. *Dermatology*, **2013**, 226, 365-70	4.4	18
56	Desmoglein-1/Erbin interaction suppresses ERK activation to support epidermal differentiation. *Journal of Clinical Investigation*, **2013**, 123, 1556-70	15.9	97
55	Recommended strategies for epidermolysis bullosa management in romania. *M dica*, **2013**, 8, 200-5		2
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