Eli Sprecher

List of Publications by Citations

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	Revised nomenclature and classification of inherited ichthyoses: results of the First Ichthyosis Consensus Conference in Sorø 2009. *Journal of the American Academy of Dermatology*, **2010**, 63, 607-41	4.5	456
219	Mutations in GALNT3, encoding a protein involved in O-linked glycosylation, cause familial tumoral calcinosis. *Nature Genetics*, **2004**, 36, 579-81	36.3	448
218	Mutations in ABCA12 underlie the severe congenital skin disease harlequin ichthyosis. *American Journal of Human Genetics*, **2005**, 76, 794-803	11	244
217	Desmoglein 1 deficiency results in severe dermatitis, multiple allergies and metabolic wasting. *Nature Genetics*, **2013**, 45, 1244-1248	36.3	217
216	Consensus reclassification of inherited epidermolysis bullosa and other disorders with skin fragility. *British Journal of Dermatology*, **2020**, 183, 614-627	4	179
215	Familial pityriasis rubra pilaris is caused by mutations in CARD14. *American Journal of Human Genetics*, **2012**, 91, 163-70	11	164
214	Hypotrichosis with juvenile macular dystrophy is caused by a mutation in CDH3, encoding P-cadherin. *Nature Genetics*, **2001**, 29, 134-6	36.3	141
213	A mutation in SNAP29, coding for a SNARE protein involved in intracellular trafficking, causes a novel neurocutaneous syndrome characterized by cerebral dysgenesis, neuropathy, ichthyosis, and palmoplantar keratoderma. *American Journal of Human Genetics*, **2005**, 77, 242-51	11	139
212	The spectrum of pathogenic mutations in SPINK5 in 19 families with Netherton syndrome: implications for mutation detection and first case of prenatal diagnosis. *Journal of Investigative Dermatology*, **2001**, 117, 179-87	4.3	114
211	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
210	Desmoglein-1/Erbin interaction suppresses ERK activation to support epidermal differentiation. *Journal of Clinical Investigation*, **2013**, 123, 1556-70	15.9	97
209	Naegeli-Franceschetti-Jadassohn syndrome and dermatopathia pigmentosa reticularis: two allelic ectodermal dysplasias caused by dominant mutations in KRT14. *American Journal of Human Genetics*, **2006**, 79, 724-30	11	94
208	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
207	Identification of a recurrent mutation in GALNT3 demonstrates that hyperostosis-hyperphosphatemia syndrome and familial tumoral calcinosis are allelic disorders. *Journal of Molecular Medicine*, **2005**, 83, 33-8	5.5	90
206	Evidence for novel functions of the keratin tail emerging from a mutation causing ichthyosis hystrix. *Journal of Investigative Dermatology*, **2001**, 116, 511-9	4.3	88
205	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
204	Molecular analysis of 250 patients with autosomal recessive congenital ichthyosis: evidence for mutation hotspots in ALOXE3 and allelic heterogeneity in ALOX12B. *Journal of Investigative Dermatology*, **2009**, 129, 1421-8	4.3	77

203	Stabilizing mutations of KLHL24 ubiquitin ligase cause loss of keratin 14 and human skin fragility. *Nature Genetics*, **2016**, 48, 1508-1516	36.3	73
202	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
201	Population-specific association between a polymorphic variant in ST18, encoding a pro-apoptotic molecule, and pemphigus vulgaris. *Journal of Investigative Dermatology*, **2012**, 132, 1798-805	4.3	71
200	Mechanisms Causing Loss of Keratinocyte Cohesion in Pemphigus. *Journal of Investigative Dermatology*, **2018**, 138, 32-37	4.3	70
199	Hyperphosphatemic familial tumoral calcinosis caused by a mutation in GALNT3 in a European kindred. *Journal of Human Genetics*, **2006**, 51, 487-490	4.3	66
198	P-cadherin regulates human hair growth and cycling via canonical Wnt signaling and transforming growth factor-β. *Journal of Investigative Dermatology*, **2012**, 132, 2332-2341	4.3	64
197	Familial tumoral calcinosis: from characterization of a rare phenotype to the pathogenesis of ectopic calcification. *Journal of Investigative Dermatology*, **2010**, 130, 652-60	4.3	61
196	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
195	CEDNIK syndrome results from loss-of-function mutations in SNAP29. *British Journal of Dermatology*, **2011**, 164, 610-6	4	59
194	Alopecia, neurological defects, and endocrinopathy syndrome caused by decreased expression of RBM28, a nucleolar protein associated with ribosome biogenesis. *American Journal of Human Genetics*, **2008**, 82, 1114-21	11	57
193	Loss of SNAP29 impairs endocytic recycling and cell motility. *PLoS ONE*, **2010**, 5, e9759	3.7	53
192	A mutation in LIPN, encoding epidermal lipase N, causes a late-onset form of autosomal-recessive congenital ichthyosis. *American Journal of Human Genetics*, **2011**, 88, 482-7	11	51
191	Galli-Galli disease is an acantholytic variant of Dowling-Degos disease. *British Journal of Dermatology*, **2007**, 156, 572-4	4	49
190	Variant in Central Centrifugal Cicatricial Alopecia. *New England Journal of Medicine*, **2019**, 380, 833-841	59.2	49
189	Monopathogenic vs multipathogenic explanations of pemphigus pathophysiology. *Experimental Dermatology*, **2016**, 25, 839-846	4	48
188	The molecular genetic analysis of the expanding pachyonychia congenita case collection. *British Journal of Dermatology*, **2014**, 171, 343-55	4	48
187	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
186	Epidermolysis bullosa simplex in Israel: clinical and genetic features. *Archives of Dermatology*, **2003**, 139, 498-505		48

185	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
184	Homozygous splice site mutations in PKP1 result in loss of epidermal plakophilin 1 expression and underlie ectodermal dysplasia/skin fragility syndrome in two consanguineous families. *Journal of Investigative Dermatology*, **2004**, 122, 647-51	4.3	43
183	Inflammatory peeling skin syndrome caused by a mutation in CDSN encoding corneodesmosin. *Journal of Investigative Dermatology*, **2011**, 131, 779-81	4.3	42
182	Molecular epidemiology of hereditary epidermolysis bullosa in a Middle Eastern population. *Journal of Investigative Dermatology*, **2006**, 126, 777-81	4.3	42
181	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
180	Peeling off the genetics of atopic dermatitis-like congenital disorders. *Journal of Allergy and Clinical Immunology*, **2014**, 134, 808-15	11.5	39
179	A mutation in a skin-specific isoform of SMARCAD1 causes autosomal-dominant adermatoglyphia. *American Journal of Human Genetics*, **2011**, 89, 302-7	11	39
178	Identification of a Functional Risk Variant for Pemphigus Vulgaris in the ST18 Gene. *PLoS Genetics*, **2016**, 12, e1006008	6	38
177	The Genetics of Pemphigus Vulgaris. *Frontiers in Medicine*, **2018**, 5, 226	4.9	36
176	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
175	Epidermolysis bullosa simplex. *Dermatologic Clinics*, **2010**, 28, 23-32	4.2	34
174	Epidermolysis bullosa simplex with mottled pigmentation resulting from a recurrent mutation in KRT14. *Journal of Investigative Dermatology*, **2006**, 126, 1654-7	4.3	34
173	Epidermolytic hyperkeratosis and epidermolysis bullosa simplex caused by frameshift mutations altering the v2 tail domains of keratin 1 and keratin 5. *Journal of Investigative Dermatology*, **2003**, 120, 623-6	4.3	33
172	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
171	Olmsted syndrome caused by a homozygous recessive mutation in TRPV3. *Journal of Investigative Dermatology*, **2014**, 134, 1752-1754	4.3	32
170	Atrichia with papular lesions maps to 8p in the region containing the human hairless gene. *American Journal of Medical Genetics Part A*, **1998**, 80, 546-50		31
169	Best treatment practices for pachyonychia congenita. *Journal of the European Academy of Dermatology and Venereology*, **2014**, 28, 279-85	4.6	30
168	Genetic hair and nail disorders. *Clinics in Dermatology*, **2005**, 23, 47-55	3	30

167	Cole Disease Results from Mutations in ENPP1. *American Journal of Human Genetics*, **2013**, 93, 752-7	11	28
166	Disadhesion of epidermal keratinocytes: a histologic clue to palmoplantar keratodermas caused by DSG1 mutations. *Journal of the American Academy of Dermatology*, **2010**, 62, 107-113	4.5	28
165	The immigration delay disease: adermatoglyphia-inherited absence of epidermal ridges. *Journal of the American Academy of Dermatology*, **2011**, 64, 974-80	4.5	27
164	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
163	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
162	Rapid detection of homozygous mutations in congenital recessive ichthyosis. *Archives of Dermatological Research*, **2008**, 300, 81-5	3.3	26
161	Homozygosity mapping as a screening tool for the molecular diagnosis of hereditary skin diseases in consanguineous populations. *Journal of the American Academy of Dermatology*, **2006**, 55, 393-401	4.5	26
160	Diffuse nonepidermolytic palmoplantar keratoderma caused by a recurrent nonsense mutation in DSG1. *Archives of Dermatology*, **2005**, 141, 625-8		26
159	Tinea capitis outbreak among paediatric refugee population, an evolving healthcare challenge. *Mycoses*, **2016**, 59, 553-7	5.2	26
158	Inherited desmosomal disorders. *Cell and Tissue Research*, **2015**, 360, 457-75	4.2	25
157	The expanding spectrum of IgA pemphigus: a case report and review of the literature. *British Journal of Dermatology*, **2014**, 171, 650-6	4	25
156	Inflammatory peeling skin syndrome caused a novel mutation in CDSN. *Archives of Dermatological Research*, **2012**, 304, 251-5	3.3	25
155	Treatment of epidermolysis bullosa pruriginosa-associated pruritus with dupilumab. *British Journal of Dermatology*, **2020**, 182, 1495-1497	4	25
154	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
153	Non-syndromic autosomal recessive congenital ichthyosis in the Israeli population. *Clinical and Experimental Dermatology*, **2013**, 38, 911-6	1.8	24
152	Abca12-mediated lipid transport and Snap29-dependent trafficking of lamellar granules are crucial for epidermal morphogenesis in a zebrafish model of ichthyosis. *DMM Disease Models and Mechanisms*, **2011**, 4, 777-85	4.1	23
151	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
150	Topobiology of human pigmentation: P-cadherin selectively stimulates hair follicle melanogenesis. *Journal of Investigative Dermatology*, **2013**, 133, 1591-600	4.3	22

149	Novel mutations in DSG1 causing striate palmoplantar keratoderma. *Clinical and Experimental Dermatology*, **2009**, 34, 224-8	1.8	22
148	Segmental basal cell naevus syndrome caused by an activating mutation in smoothened. *British Journal of Dermatology*, **2016**, 175, 178-81	4	22
147	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
146	Analysis of CARD14 Polymorphisms in Pityriasis Rubra Pilaris: Activation of NF-□B. *Journal of Investigative Dermatology*, **2015**, 135, 1905-1908	4.3	20
145	Mutations in TSPEAR, Encoding a Regulator of Notch Signaling, Affect Tooth and Hair Follicle Morphogenesis. *PLoS Genetics*, **2016**, 12, e1006369	6	20
144	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
143	Identification of a novel locus associated with congenital recessive ichthyosis on 12p11.2-q13. *Journal of Investigative Dermatology*, **2005**, 125, 456-62	4.3	19
142	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
141	Unveiling the roots of monogenic genodermatoses: genotrichoses as a paradigm. *Journal of Investigative Dermatology*, **2012**, 132, 906-14	4.3	18
140	ANE syndrome caused by mutated RBM28 gene: a novel etiology of combined pituitary hormone deficiency. *European Journal of Endocrinology*, **2010**, 162, 1021-5	6.5	17
139	Chronic pain in pachyonychia congenita: evidence for neuropathic origin. *British Journal of Dermatology*, **2018**, 179, 154-162	4	16
138	Deleterious mutations in SPINK5 in a patient with congenital ichthyosiform erythroderma: molecular testing as a helpful diagnostic tool for Netherton syndrome. *Clinical and Experimental Dermatology*, **2004**, 29, 513-7	1.8	16
137	Revisiting pachyonychia congenita: a case-cohort study of 815 patients. *British Journal of Dermatology*, **2020**, 182, 738-746	4	16
136	Assessment of the effectiveness of topical propranolol 4% gel for infantile hemangiomas. *International Journal of Dermatology*, **2017**, 56, 148-153	1.7	15
135	A distinct cutaneous microbiota profile in autoimmune bullous disease patients. *Experimental Dermatology*, **2017**, 26, 1221-1227	4	15
134	SAM syndrome is characterized by extensive phenotypic heterogeneity. *Experimental Dermatology*, **2018**, 27, 787-790	4	15
133	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
132	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

131	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
130	RBM28, a protein deficient in ANE syndrome, regulates hair follicle growth via miR-203 and p63. *Experimental Dermatology*, **2015**, 24, 618-22	4	13
129	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
128	Digenic inheritance in epidermolysis bullosa simplex. *Journal of Investigative Dermatology*, **2012**, 132, 2852-4	4.3	13
127	Paraneoplastic pityriasis rubra pilaris: case report and literature review. *Clinical and Experimental Dermatology*, **2017**, 42, 54-57	1.8	12
126	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
125	ST18 Enhances PV-IgG-Induced Loss of Keratinocyte Cohesion in Parallel to Increased ERK Activation. *Frontiers in Immunology*, **2019**, 10, 770	8.4	11
124	Topical pimecrolimus for paediatric cutaneous mastocytosis. *Clinical and Experimental Dermatology*, **2018**, 43, 559-565	1.8	11
123	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
122	IGFBP7 as a potential therapeutic target in Psoriasis. *Journal of Investigative Dermatology*, **2011**, 131, 1767-70	4.3	11
121	Tumoral calcinosis: new insights for the rheumatologist into a familial crystal deposition disease. *Current Rheumatology Reports*, **2007**, 9, 237-42	4.9	11
120	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
119	Novel TGM5 mutations in acral peeling skin syndrome. *Experimental Dermatology*, **2015**, 24, 285-9	4	10
118	Treatment of hereditary hypotrichosis simplex of the scalp with topical gentamicin. *British Journal of Dermatology*, **2020**, 183, 114-120	4	10
117	Papillon-Lef□re syndrome: report of six patients and identification of a novel mutation. *International Journal of Dermatology*, **2016**, 55, 898-902	1.7	9
116	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
115	Pachyonychia congenita cornered: report on the 11th Annual International Pachyonychia Congenita Consortium Meeting. *British Journal of Dermatology*, **2014**, 171, 974-7	4	9
114	Immunogenicity of a BNT162b2 vaccine booster in health-care workers. *Lancet Microbe, The*, **2021**, 2, e650	22.2	9

113	The Molecular Revolution in Cutaneous Biology: Era of Next-Generation Sequencing. *Journal of Investigative Dermatology*, **2017**, 137, e79-e82	4.3	8
112	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
111	Griseofulvin vs terbinafine for paediatric tinea capitis: When and for how long. *Mycoses*, **2019**, 62, 949-953	3.2	8
110	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
109	Molecular analysis of a series of Israeli families with Com Netherton syndrome. *Dermatology*, **2014**, 228, 183-8	4.4	8
108	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
107	NB-UVB phototherapy for generalized granuloma annulare. *Dermatologic Therapy*, **2016**, 29, 152-4	2.2	8
106	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
105	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
104	Striate palmoplantar keratoderma resulting from a missense mutation in DSG1. *British Journal of Dermatology*, **2018**, 179, 755-757	4	7
103	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
102	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
101	Semidominant inheritance in epidermolytic ichthyosis. *Journal of Investigative Dermatology*, **2013**, 133, 2626-2628	4.3	7
100	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
99	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
98	Loss-of-function variants in C3ORF52 result in localized autosomal recessive hypotrichosis. *Genetics in Medicine*, **2020**, 22, 1227-1234	8.1	6
97	Mucous membrane pemphigoid-otorhinolaryngological manifestations: a retrospective cohort study. *European Archives of Oto-Rhino-Laryngology*, **2020**, 277, 939-945	3.5	6
96	Childhood Pemphigus Foliaceus with Exclusive Immunoglobulin G Autoantibodies to Desmocollins. *Pediatric Dermatology*, **2016**, 33, e10-3	1.9	6

95	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
94	SVEP1 plays a crucial role in epidermal differentiation. *Experimental Dermatology*, **2017**, 26, 423-430	4	6
93	Reduced folate carrier (RFC-1) gene expression in normal and psoriatic skin. *Archives of Dermatological Research*, **1998**, 290, 656-60	3.3	6
92	Non-keratinocyte SNAP29 influences epidermal differentiation and hair follicle formation in mice. *Experimental Dermatology*, **2016**, 25, 647-9	4	6
91	Effectiveness of topical propranolol 4% gel in the treatment of pyogenic granuloma in children. *Journal of Dermatology*, **2019**, 46, 245-248	1.6	6
90	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
89	The pathogenesis of melasma and implications for treatment. *Journal of Cosmetic Dermatology*, **2021**, 20, 3432-3445	2.5	6
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	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
86	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
85	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
84	A refractory, cutaneous, subepidermal bullous disease. *Clinical and Experimental Dermatology*, **2016** , 41, 573-5	1.8	5
83	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
82	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
81	Occupational mycosis fungoides - a case series. *International Journal of Dermatology*, **2017**, 56, 733-737	1.7	4
80	Report of the 13th Annual International Pachyonychia Congenita Consortium Symposium. *British Journal of Dermatology*, **2017**, 176, 1144-1147	4	4
79	PLACK syndrome shows remarkable phenotypic homogeneity. *Clinical and Experimental Dermatology*, **2019**, 44, 580-583	1.8	4
78	Glutathione S-transferase polymorphisms in patients with photosensitive and non-photosensitive drug eruptions. *Photodermatology Photoimmunology and Photomedicine*, **2019**, 35, 214-220	2.4	4

77	Intense focused ultrasound for neck and lower face skin tightening a prospective study. *Journal of Cosmetic Dermatology*, **2020**, 19, 850-854	2.5	4
76	Loss-of-Function Variants in SERPINA12 Underlie Autosomal Recessive Palmoplantar Keratoderma. *Journal of Investigative Dermatology*, **2020**, 140, 2178-2187	4.3	4
75	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
74	Punctate palmoplantar keratoderma: an unusual mutation causing an unusual phenotype. *British Journal of Dermatology*, **2018**, 178, 1455-1457	4	4
73	A novel homozygous deletion in EXPH5 causes a skin fragility phenotype. *Clinical and Experimental Dermatology*, **2016**, 41, 915-918	1.8	4
72	Bullous pemphigoid and diabetes mellitus: Are we missing the larger picture?. *Journal of the American Academy of Dermatology*, **2018**, 79, e27	4.5	4
71	Translational implications of Th17-skewed inflammation due to genetic deficiency of a cadherin stress sensor.. *Journal of Clinical Investigation*, **2021**,	15.9	4
70	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
69	An enhanced transcutaneous delivery of botulinum toxin for the treatment of Hailey-Hailey disease. *Dermatologic Therapy*, **2020**, 33, e13184	2.2	4
68	Grover disease and bullous pemphigoid: a clinicopathological study of six cases. *Clinical and Experimental Dermatology*, **2019**, 44, 524-527	1.8	4
67	Anterior Scleritis Associated with Pemphigus Vulgaris. *Ocular Immunology and Inflammation*, **2019**, 27, 497-498	2.8	4
66	Comorbidities in patients with palmoplantar plaque psoriasis. *Journal of the American Academy of Dermatology*, **2021**, 84, 639-643	4.5	4
65	Clinical efficacy of fecal microbial transplantation treatment in adults with moderate-to-severe atopic dermatitis.. *Immunity, Inflammation and Disease*, **2021**,	2.4	4
64	ARCI7 Revisited and Repositioned. *Journal of Investigative Dermatology*, **2017**, 137, 970-972	4.3	3
63	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
62	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
61	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
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