

Alon Peled

List of Publications by Year in descending order

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Version: 2024-02-01

27
papers

487
citations

933264

10
h-index

713332

21
g-index

28
all docs

28
docs citations

28
times ranked

843
citing authors

#	ARTICLE	IF	CITATIONS
1	Variant <i>PADI3</i> in Central Centrifugal Cicatricial Alopecia. <i>New England Journal of Medicine</i> , 2019, 380, 833-841.	13.9	102
2	Loss-of-function mutations in caspase recruitment domain-containing protein 14 (CARD14) are associated with a severe variant of atopic dermatitis. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 173-181.e10.	1.5	60
3	Identification of a Functional Risk Variant for Pemphigus Vulgaris in the ST18 Gene. <i>PLoS Genetics</i> , 2016, 12, e1006008.	1.5	53
4	A phenotype combining hidradenitis suppurativa with Dowling-Degos disease caused by a founder mutation in <i>PSENEN</i> . <i>British Journal of Dermatology</i> , 2018, 178, 502-508.	1.4	48
5	Filaggrin 2 Deficiency Results in Abnormal Cell-Cell Adhesion in the Cornified Cell Layers and Causes Peeling Skin Syndrome Type A. <i>Journal of Investigative Dermatology</i> , 2018, 138, 1736-1743.	0.3	37
6	Mutations in TSPEAR, Encoding a Regulator of Notch Signaling, Affect Tooth and Hair Follicle Morphogenesis. <i>PLoS Genetics</i> , 2016, 12, e1006369.	1.5	32
7	Calpain 12 Function Revealed through the Study of an Atypical Case of Autosomal Recessive Congenital Ichthyosis. <i>Journal of Investigative Dermatology</i> , 2017, 137, 385-393.	0.3	19
8	Treatment of hereditary hypotrichosis simplex of the scalp with topical gentamicin. <i>British Journal of Dermatology</i> , 2020, 183, 114-120.	1.4	19
9	The Role of Desmoglein 1 in Gap Junction Turnover Revealed through the Study of SAM Syndrome. <i>Journal of Investigative Dermatology</i> , 2020, 140, 556-567.e9.	0.3	17
10	Loss-of-Function Variants in SERPINA12 Underlie Autosomal Recessive Palmoplantar Keratoderma. <i>Journal of Investigative Dermatology</i> , 2020, 140, 2178-2187.	0.3	14
11	Recessive epidermolytic ichthyosis results from loss of keratin 10 expression, regardless of the mutation location. <i>Clinical and Experimental Dermatology</i> , 2018, 43, 187-190.	0.6	10
12	Hypertension and childhood migration. <i>Journal of Hypertension</i> , 2019, 37, 702-709.	0.3	10
13	Immigration to Israel during childhood is associated with diabetes at adolescence: a study of 2.7 million adolescents. <i>Diabetologia</i> , 2017, 60, 2226-2230.	2.9	9
14	Myopia and Childhood Migration. <i>Ophthalmology</i> , 2020, 127, 713-723.	2.5	7
15	ST18 affects cell-cell adhesion in pemphigus vulgaris in a tumour necrosis factor-dependent fashion*. <i>British Journal of Dermatology</i> , 2021, 184, 1153-1160.	1.4	7
16	Coexistence of pachyonychia congenita and hidradenitis suppurativa: more than a coincidence. <i>British Journal of Dermatology</i> , 2022, 187, 392-400.	1.4	7
17	Somatic Mosaicism for a <i>GJB2</i> Mutation Results in a Patterned Form of Spiny Hyperkeratosis without Eccrine Involvement. <i>Pediatric Dermatology</i> , 2016, 33, 322-326.	0.5	5
18	Palmoplantar keratoderma caused by a missense variant in <i>CTSB</i> encoding cathepsin B. <i>Clinical and Experimental Dermatology</i> , 2021, 46, 103-108.	0.6	5

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19	Autosomal dominant cutis laxa resulting from an intronic mutation in <i>ELN</i> . <i>Experimental Dermatology</i> , 2015, 24, 885-887.	1.4	4
20	Diffuse Facial Hyperpigmentation as a Presenting Sign of Lupus Erythematosus: Three Cases and Review of the Literature. <i>Case Reports in Dermatology</i> , 2021, 13, 263-270.	0.3	4
21	Laryngeal mucous membrane pemphigoid serves as a prognostic factor for poor response to treatment with rituximab. <i>Clinical and Experimental Dermatology</i> , 2021, 46, 915-919.	0.6	3
22	Myopia and Early-Onset Type 2 Diabetes: A Nationwide Cohort Study. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, e663-e671.	1.8	3
23	Linear accelerator stereotactic radiosurgery can modulate the clinical course of Hemangioblastoma: Case series and review of the literature. <i>Journal of Clinical Neuroscience</i> , 2020, 82, 162-165.	0.8	3
24	Loss-of-function variants in KLF4 underlie autosomal dominant palmoplantar keratoderma. <i>Genetics in Medicine</i> , 2022, 24, 1085-1095.	1.1	3
25	Phenotypic suppression of acral peeling skin syndrome in a patient with autosomal recessive congenital ichthyosis. <i>Experimental Dermatology</i> , 2020, 29, 742-748.	1.4	2
26	Postoperative supine position after primary Descemet-stripping automated endothelial keratoplasty reduces graft detachment rate. <i>Canadian Journal of Ophthalmology</i> , 2022, 57, 147-153.	0.4	2
27	Molecular epidemiology of pachyonychia congenita in the Israeli population. <i>Clinical and Experimental Dermatology</i> , 2021, 46, 663-668.	0.6	1