Alon Peled

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/554276/publications.pdf

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27	487	933264	713332
papers	citations	10 h-index	21 g-index

28	28	28	843
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Postoperative supine position after primary Descemet-stripping automated endothelial keratoplasty reduces graft detachment rate. Canadian Journal of Ophthalmology, 2022, 57, 147-153.	0.4	2
2	Myopia and Early-Onset Type 2 Diabetes: A Nationwide Cohort Study. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e663-e671.	1.8	3
3	Loss-of-function variants in KLF4 underlie autosomal dominant palmoplantar keratoderma. Genetics in Medicine, 2022, 24, 1085-1095.	1.1	3
4	Coexistence of pachyonychia congenita and hidradenitis suppurativa: more than a coincidence. British Journal of Dermatology, 2022, 187, 392-400.	1.4	7
5	Palmoplantar keratoderma caused by a missense variant in CTSB encoding cathepsin B. Clinical and Experimental Dermatology, 2021, 46, 103-108.	0.6	5
6	Molecular epidemiology of pachyonychia congenita in the Israeli population. Clinical and Experimental Dermatology, 2021, 46, 663-668.	0.6	1
7	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.	1.4	7
8	Laryngeal mucous membrane pemphigoid serves as a prognostic factor for poor response to treatment with rituximab. Clinical and Experimental Dermatology, 2021, 46, 915-919.	0.6	3
9	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.	0.3	4
10	The Role of Desmoglein 1 in Gap Junction Turnover Revealed through the Study of SAMÂSyndrome. Journal of Investigative Dermatology, 2020, 140, 556-567.e9.	0.3	17
11	Myopia and Childhood Migration. Ophthalmology, 2020, 127, 713-723.	2.5	7
12	Treatment of hereditary hypotrichosis simplex of the scalp with topical gentamicin. British Journal of Dermatology, 2020, 183, 114-120.	1.4	19
13	Phenotypic suppression of acral peeling skin syndrome in a patient with autosomal recessive congenital ichthyosis. Experimental Dermatology, 2020, 29, 742-748.	1.4	2
14	Loss-of-Function Variants in SERPINA12 Underlie Autosomal Recessive Palmoplantar Keratoderma. Journal of Investigative Dermatology, 2020, 140, 2178-2187.	0.3	14
15	Linear accelerator stereotactic radiosurgery can modulate the clinical course of Hemangioblastoma: Case series and review of the literature. Journal of Clinical Neuroscience, 2020, 82, 162-165.	0.8	3
16	Variant PADI3 in Central Centrifugal Cicatricial Alopecia. New England Journal of Medicine, 2019, 380, 833-841.	13.9	102
17	Hypertension and childhood migration. Journal of Hypertension, 2019, 37, 702-709.	0.3	10
18	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.	1.5	60

#	Article	IF	Citations
19	Recessive epidermolytic ichthyosis results from loss of keratin 10 expression, regardless of the mutation location. Clinical and Experimental Dermatology, 2018, 43, 187-190.	0.6	10
20	A phenotype combining hidradenitis suppurativa with Dowling-Degos disease caused by a founder mutation in PSENEN. British Journal of Dermatology, 2018, 178, 502-508.	1.4	48
21	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.	0.3	37
22	Calpain 12 Function Revealed through the Study of an Atypical Case of Autosomal Recessive Congenital Ichthyosis. Journal of Investigative Dermatology, 2017, 137, 385-393.	0.3	19
23	Immigration to Israel during childhood is associated with diabetes at adolescence: a study of 2.7 million adolescents. Diabetologia, 2017, 60, 2226-2230.	2.9	9
24	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.	0.5	5
25	Identification of a Functional Risk Variant for Pemphigus Vulgaris in the ST18 Gene. PLoS Genetics, 2016, 12, e1006008.	1.5	53
26	Mutations in TSPEAR, Encoding a Regulator of Notch Signaling, Affect Tooth and Hair Follicle Morphogenesis. PLoS Genetics, 2016, 12, e1006369.	1.5	32
27	Autosomalâ€dominant cutis laxa resulting from an intronic mutation in <scp>ELN</scp>. Experimental Dermatology, 2015, 24, 885-887.	1.4	4