Janan Mohamad

List of Publications by Year
in descending order

13	298	1162889 8	1058333 14
papers	citations	h-index	g-index

14	14	14	445
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Loss-of-function variants in KLF4 underlie autosomal dominant palmoplantar keratoderma. Genetics in Medicine, 2022, 24, 1085-1095.	1.1	3
2	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.	1.6	1
3	Epidermolytic epidermal nevus caused by a somatic mutation in KRT2. Pediatric Dermatology, 2021, 38, 538-540.	0.5	1
4	Molecular epidemiology of nonâ€syndromic autosomal recessive congenital ichthyosis in a Middleâ€Eastern population. Experimental Dermatology, 2021, 30, 1290-1297.	1.4	10
5	Phenotypic suppression of acral peeling skin syndrome in a patient with autosomal recessive congenital ichthyosis. Experimental Dermatology, 2020, 29, 742-748.	1.4	2
6	Loss-of-function variants in C3ORF52 result in localized autosomal recessive hypotrichosis. Genetics in Medicine, 2020, 22, 1227-1234.	1.1	12
7	Loss-of-Function Variants in SERPINA12 Underlie Autosomal Recessive Palmoplantar Keratoderma. Journal of Investigative Dermatology, 2020, 140, 2178-2187.	0.3	14
8	Variant PADI3 in Central Centrifugal Cicatricial Alopecia. New England Journal of Medicine, 2019, 380, 833-841.	13.9	102
9	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
10	SAM syndrome is characterized by extensive phenotypic heterogeneity. Experimental Dermatology, 2018, 27, 787-790.	1.4	22
11	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
12	ARCI7 Revisited and Repositioned. Journal of Investigative Dermatology, 2017, 137, 970-972.	0.3	6
13	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.	2.6	27