Tom Rabinowitz

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Variant <i>PADI3</i> in Central Centrifugal Cicatricial Alopecia. New England Journal of Medicine, 2019, 380, 833-841.	13.9	102
2	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
3	Bayesian-based noninvasive prenatal diagnosis of single-gene disorders. Genome Research, 2019, 29, 428-438.	2.4	31
4	Genome-wide noninvasive prenatal diagnosis of monogenic disorders: Current and future trends. Computational and Structural Biotechnology Journal, 2020, 18, 2463-2470.	1.9	22
5	Loss-of-Function Variants in SERPINA12 Underlie Autosomal Recessive Palmoplantar Keratoderma. Journal of Investigative Dermatology, 2020, 140, 2178-2187.	0.3	14
6	Molecular epidemiology of nonâ€syndromic autosomal recessive congenital ichthyosis in a Middleâ€Eastern population. Experimental Dermatology, 2021, 30, 1290-1297.	1.4	10
7	A novel intronic mutation of is a major cause of autosomal recessive retinitis pigmentosa among Caucasus Jews. Molecular Vision, 2019, 25, 155-164.	1.1	9
8	<i>PNPT1</i> , <i>MYO15A</i> , <i>PTPRQ</i> , and <i>SLC12A2</i> â€associated genetic and phenotypic heterogeneity among hearing impaired assortative mating families in Southern India. Annals of Human Genetics, 2022, 86, 1-13.	0.3	5
9	Genome-Wide Noninvasive Prenatal Diagnosis of De Novo Mutations. Methods in Molecular Biology, 2021, 2243, 249-269.	0.4	2
10	Improved noninvasive fetal variant calling using standardized benchmarking approaches. Computational and Structural Biotechnology Journal, 2021, 19, 509-517.	1.9	1
11	Genome-Wide Noninvasive Prenatal Diagnosis of SNPs and Indels. Methods in Molecular Biology, 2021, 2243, 227-248.	0.4	O