Suzanne Broadgate

List of Publications by Year
in descending order

15	269	1306789 7	1125271 13
papers	citations	h-index	g-index

15	15	15	491
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Unravelling the genetics of inherited retinal dystrophies: Past, present and future. Progress in Retinal and Eye Research, 2017, 59, 53-96.	7.3	85
2	Detailed Phenotypic and Genotypic Characterization of Bietti Crystalline Dystrophy. Ophthalmology, 2014, 121, 1174-1184.	2.5	79
3	An Overview of the Genetics of ABCA4 Retinopathies, an Evolving Story. Genes, 2021, 12, 1241.	1.0	22
4	Characterization ofCDH3-Related Congenital Hypotrichosis With Juvenile Macular Dystrophy. JAMA Ophthalmology, 2016, 134, 992.	1.4	17
5	Association of Clinical and Genetic Heterogeneity With BEST1 Sequence Variations. JAMA Ophthalmology, 2020, 138, 544.	1.4	14
6	Hypotrichosis and juvenile macular dystrophy caused by CDH3 mutation: A candidate disease for retinal gene therapy. Scientific Reports, 2016, 6, 23674.	1.6	13
7	Novel Pathogenic Sequence Variants in NR2E3 and Clinical Findings in Three Patients. Genes, 2020, 11, 1288.	1.0	8
8	Diabetic macular oedema: underâ€represented in the genetic analysis of diabetic retinopathy. Acta Ophthalmologica, 2018, 96, 1-51.	0.6	7
9	â€œGenetic and clinical findings in an ethnically diverse retinitis pigmentosa cohort associated with pathogenic variants in EYSâ€. Eye, 2021, 35, 1440-1449.	1.1	6
10	Novel homozygous splicing mutations in cause autosomal recessive retinitis pigmentosa. Molecular Vision, 2018, 24, 603-612.	1.1	6
11	Identification of rod- and cone-specific expression signatures to identify candidate genes for retinal disease. Experimental Eye Research, 2015, 132, 161-173.	1.2	5
12	Genetic and Clinical Findings in an Ethnically Diverse Cohort with Retinitis Pigmentosa Associated with Pathogenic Variants in CERKL. Genes, 2020, 11, 1497.	1.0	4
13	Targeted next generation sequencing and family survey enable correct genetic diagnosis in CRX associated macular dystrophy â€“ a case report. BMC Ophthalmology, 2021, 21, 168.	0.6	3
14	Author reply. Ophthalmology, 2015, 122, e22.	2.5	0
15	Whole genome sequencing in a Knobloch syndrome family confirms the molecular diagnosis. Ophthalmic Genetics, 2021, , 1-9.	0.5	0