VÃ-ctor Abad-Morales

List of Publications by Year
in descending order

9 papers	71 citations	1684188 5 h-index	1588992 8 g-index
9 all docs	9 docs citations	9 times ranked	135 citing authors

#	Article	IF	Citations
1	Identification of a novel homozygous ARSG mutation as the second cause of Usher syndrome type 4. American Journal of Ophthalmology Case Reports, 2020, 19, 100736.	0.7	21
2	Expanding the retinal phenotype of RP1: from retinitis pigmentosa to a novel and singular macular dystrophy. British Journal of Ophthalmology, 2020, 104, 173-181.	3.9	13
3	Expression Atlas of the Deubiquitinating Enzymes in the Adult Mouse Retina, Their Evolutionary Diversification and Phenotypic Roles. PLoS ONE, 2016, 11, e0150364.	2.5	10
4	mRNA expression analysis of the SUMO pathway genes in the adult mouse retina. Biology Open, 2015, 4, 224-232.	1.2	6
5	Characterization of the cone-rod dystrophy retinal phenotype caused by novel homozygous DRAM2 mutations. Experimental Eye Research, 2019, 187, 107752.	2.6	6
6	New Insights on the Genetic Basis Underlying SHILCA Syndrome: Characterization of the NMNAT1 Pathological Alterations Due to Compound Heterozygous Mutations and Identification of a Novel Alternative Isoform. International Journal of Molecular Sciences, 2021, 22, 2262.	4.1	6
7	Generation of an induced pluripotent stem cell line (FRIMOi007-A) derived from an incomplete achromatopsia patient carrying a novel homozygous mutation in PDE6C gene. Stem Cell Research, 2019, 40, 101569.	0.7	3
8	Generation of Best disease-derived induced pluripotent stem cell line (FRIMOi006-A) carrying a novel dominant mutation in BEST1 gene. Stem Cell Research, 2019, 40, 101570.	0.7	3
9	Coexistence of Meesmann Corneal Dystrophy and a Pseudo-Unilateral Lattice Corneal Dystrophy in a Patient With a Novel Pathogenic Variant in the Keratin K3 Gene: A Case Report. Cornea, 2021, 40, 370-372.	1.7	3