VÃ-ctor Abad-Morales

List of Publications by Year in descending order

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

#	Article	IF	CITATIONS
1	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
2	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
3	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
4	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
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	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
7	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
8	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
9	mRNA expression analysis of the SUMO pathway genes in the adult mouse retina. Biology Open, 2015, 4, 224-232.	1.2	6