

Jesús-José Ferre-Fernández

List of Publications by Year in descending order

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Version: 2024-02-01

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203
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#	ARTICLE	IF	CITATIONS
1	Hypo- and Hypermorphic FOXC1 Mutations in Dominant Glaucoma: Transactivation and Phenotypic Variability. PLoS ONE, 2015, 10, e0119272.	2.5	24
2	CPAMD8 loss-of-function underlies non-dominant congenital glaucoma with variable anterior segment dysgenesis and abnormal extracellular matrix. Human Genetics, 2020, 139, 1209-1231.	3.8	23
3	Whole-Exome Sequencing of Congenital Glaucoma Patients Reveals Hypermorphic Variants in GPATCH3, a New Gene Involved in Ocular and Craniofacial Development. Scientific Reports, 2017, 7, 46175.	3.3	22
4	Disruption of foxc1 genes in zebrafish results in dosage-dependent phenotypes overlapping Axenfeld-Rieger syndrome. Human Molecular Genetics, 2020, 29, 2723-2735.	2.9	15
5	Role of FOXC2 and PITX2 rare variants associated with mild functional alterations as modifier factors in congenital glaucoma. PLoS ONE, 2019, 14, e0211029.	2.5	10
6	Role of GUCA1C in Primary Congenital Glaucoma and in the Retina: Functional Evaluation in Zebrafish. Genes, 2020, 11, 550.	2.4	10
7	The Role of hsa-miR-548l Dysregulation as a Putative Modifier Factor for Glaucoma-Associated FOXC1 Mutations. MicroRNA (Sharjah, United Arab Emirates), 2015, 4, 50-56.	1.2	8
8	Functional characterization of eight rare missense CYP11B1 variants involved in congenital glaucoma and their association with null genotypes. Acta Ophthalmologica, 2016, 94, e555-e560.	1.1	8
9	Apparent Radiological Improvement in an Infant With Labrune Syndrome Treated With Bevacizumab. Pediatric Neurology, 2020, 112, 53-55.	2.1	7
10	Knockout of myoc Provides Evidence for the Role of Myocilin in Zebrafish Sex Determination Associated with Wnt Signalling Downregulation. Biology, 2021, 10, 98.	2.8	2
11	C-Terminal-PEDF Reduces IC50 Doses and Chemoresistant Population of CD133 and BCRP1-Positive Cancer Stem Like Cells. Journal of Analytical Oncology, 0, , .	0.1	2