A novel missense mutation in the gene for gap-junction with autosomal dominant "nuclear punctate" cataracts

Molecular Vision

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Citation Report

#	Article	IF	Citations
1	Intracellular Distribution, Assembly and Effect of Disease-associated Connexin 31 Mutants in HeLa Cells. Acta Biochimica Et Biophysica Sinica, 2005, 37, 547-554.	2.0	24
2	Two novel mutations of connexin genes in Chinese families with autosomal dominant congenital nuclear cataract. British Journal of Ophthalmology, 2005, 89, 1535-1537.	3.9	31
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5	Knock-in of Î±3 connexin prevents severe cataracts caused by an Î±8 point mutation. Journal of Cell Science, 2006, 119, 2138-2144.	2.0	34
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16	Connexin Mutants and Cataracts. Frontiers in Pharmacology, 2013, 4, 43.	3.5	90
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18	Sporadic and Familial Congenital Cataracts: Mutational Spectrum and New Diagnoses Using Nextâ€Generation Sequencing. Human Mutation, 2016, 37, 371-384.	2.5	108

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19	Inherited Congenital Cataract: A Guide to Suspect the Genetic Etiology in the Cataract Genesis. Molecular Syndromology, 2017, 8, 58-78.	0.8	58
20	â€œAnt-eggâ€•cataract revisited. Graefe's Archive for Clinical and Experimental Ophthalmology, 2017, 255, 119-125.	1.9	1
21	The connexin 46 mutant (V44M) impairs gap junction function causing congenital cataract. Journal of Genetics, 2017, 96, 969-976.	0.7	5
22	Intercellular Communication in Lens Development and Disease. , 2005, , 173-195.		2
23	Further evidence for P59L mutation in GJA3 associated with autosomal dominant congenital cataract. Indian Journal of Ophthalmology, 2016, 64, 508.	1.1	9
24	Lens Proteins and Their Molecular Biology. , 2008, , 1341-1364.		1
25	Connexins in Lens Development and Disease. , 2009, , 387-396.		1
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