

CITATION REPORT

List of articles citing

Mutations in RAB28, encoding a farnesylated small GTPase, are associated with autosomal-recessive cone-rod dystrophy

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American Journal of Human Genetics, 2013, 93, 110-7.

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#	Paper	IF	Citations
80	Genetic architecture of retinal and macular degenerative diseases: the promise and challenges of next-generation sequencing. <i>Genome Medicine</i> , 2013 , 5, 84	14.4	28
79	Whole exome sequencing reveals genetic predisposition in a large family with retinitis pigmentosa. <i>BioMed Research International</i> , 2014 , 2014, 302487	3	5
78	Membrane protein transport in photoreceptors: the function of PDE β the Proctor lecture. <i>Investigative Ophthalmology and Visual Science</i> , 2014 , 55, 8653-66		33
77	The role of primary cilia in the development and disease of the retina. <i>Organogenesis</i> , 2014 , 10, 69-85	1.7	86
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74	Prenylation defects in inherited retinal diseases. <i>Journal of Medical Genetics</i> , 2014 , 51, 143-51	5.8	19
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72	Developments in Ocular Genetics: 2013 Annual Review. <i>Asia-Pacific Journal of Ophthalmology</i> , 2014 , 3, 181-93	3.5	6
71	Disruption of the basal body protein POC1B results in autosomal-recessive cone-rod dystrophy. <i>American Journal of Human Genetics</i> , 2014 , 95, 131-42	11	47
70	Whole-exome sequencing identifies KIZ as a ciliary gene associated with autosomal-recessive rod-cone dystrophy. <i>American Journal of Human Genetics</i> , 2014 , 94, 625-33	11	42
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64	Knockdown of poc1b causes abnormal photoreceptor sensory cilium and vision impairment in zebrafish. <i>Biochemical and Biophysical Research Communications</i> , 2015 , 465, 651-7	3.4	7

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56	Whole-Organism Developmental Expression Profiling Identifies RAB-28 as a Novel Ciliary GTPase Associated with the BBSome and Intraflagellar Transport. <i>PLoS Genetics</i> , 2016 , 12, e1006469	6	33
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