An ADAM9 mutation in canine cone-rod dystrophy 3 est cone-rod dystrophy 9

Molecular Vision

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Citation Report

#	Article	IF	Citations
1	The Domestic Cat as a Large Animal Model for Characterization of Disease and Therapeutic Intervention in Hereditary Retinal Blindness. Journal of Ophthalmology, 2011, 2011, 1-8.	0.6	25
2	Characterization of feline hereditary retinal dystrophies using clinical, functional, structural and molecular genetic studies. Veterinary Ophthalmology, 2011, 14, 30-36.	0.6	7
3	Exclusion ofRPGRIP1ins44 from Primary Causal Association with Early-Onset Coneâ€“Rod Dystrophy in Dogs. , 2012, 53, 5486.		18
4	Assessment of Hereditary Retinal Degeneration in the English Springer Spaniel Dog and Disease Relationship to anRPGRIP1Mutation. Stem Cells International, 2012, 2012, 1-12.	1.2	6
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6	Copy number variation in the genomes of domestic animals. Animal Genetics, 2012, 43, 503-517.	0.6	116
7	Genetic and phenotypic variations of inherited retinal diseases in dogs: the power of within- and across-breed studies. Mammalian Genome, 2012, 23, 40-61.	1.0	78
8	Copy number variation in the domestic dog. Mammalian Genome, 2012, 23, 144-163.	1.0	55
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10	Drug and gene therapy of hereditary retinal disease in dog and cat models. Drug Discovery Today: Disease Models, 2013, 10, e215-e223.	1.2	3
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15	NovelADAM9homozygous mutation in a consanguineous Egyptian family with severe cone-rod dystrophy and cataract. British Journal of Ophthalmology, 2014, 98, 1718-1723.	2.1	7
16	The genetics of eye disorders in the dog. Canine Genetics and Epidemiology, 2014, 1, 3.	2.9	23
17	Causes and consequences of inherited cone disorders. Progress in Retinal and Eye Research, 2014, 42, 1-26.	7.3	127
18	Dog Models for Blinding Inherited Retinal Dystrophies. Human Gene Therapy Clinical Development, 2015, 26, 15-26.	3.2	50

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19	The genetics of inherited retinal disorders in dogs: implications for diagnosis and management. Veterinary Medicine: Research and Reports, 2016, 7, 41.	0.4	3
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21	Evaluation of artificial selection in Standard Poodles using whole-genome sequencing. Mammalian Genome, 2016, 27, 599-609.	1.0	13
22	Animal modelling for inherited central vision loss. Journal of Pathology, 2016, 238, 300-310.	2.1	50
23	Novel retinopathy in related Gordon setters: a clinical, behavioral, electrophysiological, and genetic investigation. Veterinary Ophthalmology, 2016, 19, 398-408.	0.6	5
24	Mapping of Canine Models of Inherited Retinal Diseases. Advances in Experimental Medicine and Biology, 2018, 1074, 257-264.	0.8	2
26	Peripheral Cone Dystrophy: Expanded Clinical Spectrum, Multimodal and Ultrawide-Field Imaging, and Genomic Analysis. Journal of Ophthalmology, 2018, 2018, 1-13.	0.6	3
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36	Dog Models for Blinding Inherited Retinal Degenerations. Human Gene Therapy Clinical Development, 0, , 150127063140004.	3.2	0
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