Maria Kaukonen

List of Publications by Year
in descending order

12	248	1307594 7	1125743 13
papers	citations	h-index	g-index

13	13	13	407
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A missense variant in IFT122 associated with a canine model of retinitis pigmentosa. Human Genetics, 2021, 140, 1569-1579.	3.8	4
2	Clinical and Genetic Findings in 28 American Cocker Spaniels with Aural Ceruminous Gland Hyperplasia and Ectasia. Journal of Comparative Pathology, 2021, 185, 30-44.	0.4	2
3	A Missense Variant in the Bardet-Biedl Syndrome 2 Gene (BBS2) Leads to a Novel Syndromic Retinal Degeneration in the Shetland Sheepdog. Genes, 2021, 12, 1771.	2.4	4
4	A putative silencer variant in a spontaneous canine model of retinitis pigmentosa. PLoS Genetics, 2020, 16, e1008659.	3.5	9
5	Formal commentary. PLoS Genetics, 2020, 16, e1009059.	3.5	2
6	Whole Genome Sequencing of Giant Schnauzer Dogs with Progressive Retinal Atrophy Establishes NECAP1 as a Novel Candidate Gene for Retinal Degeneration. Genes, 2019, 10, 385.	2.4	6
7	Skin microbiota and allergic symptoms associate with exposure to environmental microbes. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, 4897-4902.	7.1	51
8	Maternal Inheritance of a Recessive RBP4 Defect in Canine Congenital Eye Disease. Cell Reports, 2018, 23, 2643-2652.	6.4	17
9	Frequency and distribution of 152 genetic disease variants in over 100,000 mixed breed and purebred dogs. PLoS Genetics, 2018, 14, e1007361.	3.5	62
10	An intronic LINE-1 insertion in MERTK is strongly associated with retinopathy in Swedish Vallhund dogs. PLoS ONE, 2017, 12, e0183021.	2.5	10
11	Genetic Panel Screening of Nearly 100 Mutations Reveals New Insights into the Breed Distribution of Risk Variants for Canine Hereditary Disorders. PLoS ONE, 2016, 11, e0161005.	2.5	43
12	A Novel Missense Mutation in ADAMTS10 in Norwegian Elkhound Primary Glaucoma. PLoS ONE, 2014, 9, e111941.	2.5	34