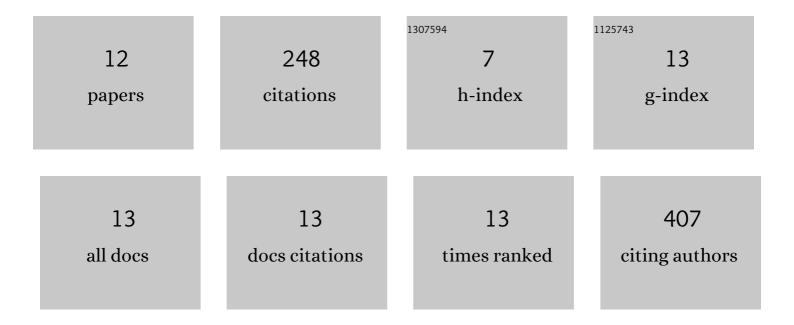
Maria Kaukonen

List of Publications by Year in descending order

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| # | 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 |