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List of Publications by Year in descending order

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Version: 2024-02-01

46
papers

1,310
citations

623188

14
h-index

414034

32
g-index

49
all docs

49
docs citations

49
times ranked

2041
citing authors

#	ARTICLE	IF	CITATIONS
1	Validation of a Chromosome 14 Risk Haplotype for Idiopathic Epilepsy in the Belgian Shepherd Dog Found to Be Associated with an Insertion in the RAPGEF5 Gene. <i>Genes</i> , 2022, 13, 1124.	1.0	7
2	An across-breed validation study of 46 genetic markers in canine hip dysplasia. <i>BMC Genomics</i> , 2021, 22, 68.	1.2	7
3	Intronic variant in POU1F1 associated with canine pituitary dwarfism. <i>Human Genetics</i> , 2021, 140, 1553-1562.	1.8	4
4	Canine DVL2 variant contributes to brachycephalic phenotype and caudal vertebral anomalies. <i>Human Genetics</i> , 2021, 140, 1535-1545.	1.8	9
5	A hypomyelinating leukodystrophy in German Shepherd dogs. <i>Journal of Veterinary Internal Medicine</i> , 2021, 35, 1455-1465.	0.6	4
6	In-frame deletion in canine PITRM1 is associated with a severe early-onset epilepsy, mitochondrial dysfunction and neurodegeneration. <i>Human Genetics</i> , 2021, 140, 1593-1609.	1.8	9
7	Missense variant in LOXHD1 is associated with canine nonsyndromic hearing loss. <i>Human Genetics</i> , 2021, 140, 1611-1618.	1.8	4
8	Clinical and Genetic Findings in 28 American Cocker Spaniels with Aural Ceruminous Gland Hyperplasia and Ectasia. <i>Journal of Comparative Pathology</i> , 2021, 185, 30-44.	0.1	2
9	Improving the resolution of canine genome-wide association studies using genotype imputation: A study of two breeds. <i>Animal Genetics</i> , 2021, 52, 703-713.	0.6	5
10	Whole Genome Sequencing Reveals Multiple Linked Genetic Variants on Canine Chromosome 12 Associated with Risk for Symmetrical Lupoid Onychodystrophy (SLO) in the Bearded Collie. <i>Genes</i> , 2021, 12, 1265.	1.0	2
11	Dog colour patterns explained by modular promoters of ancient canid origin. <i>Nature Ecology and Evolution</i> , 2021, 5, 1415-1423.	3.4	24
12	Genetic characterization of Addison's disease in Bearded Collies. <i>BMC Genomics</i> , 2020, 21, 833.	1.2	4
13	A novel genomic region on chromosome 11 associated with fearfulness in dogs. <i>Translational Psychiatry</i> , 2020, 10, 169.	2.4	10
14	Recessive missense LAMP3 variant associated with defect in lamellar body biogenesis and fatal neonatal interstitial lung disease in dogs. <i>PLoS Genetics</i> , 2020, 16, e1008651.	1.5	8
15	A putative silencer variant in a spontaneous canine model of retinitis pigmentosa. <i>PLoS Genetics</i> , 2020, 16, e1008659.	1.5	9
16	webGQT: A Shiny Server for Genotype Query Tools for Model-Based Variant Filtering. <i>Frontiers in Genetics</i> , 2020, 11, 152.	1.1	10
17	Whole Genome Sequencing Indicates Heterogeneity of Hyperostotic Disorders in Dogs. <i>Genes</i> , 2020, 11, 163.	1.0	4
18	A Missense Variant Affecting the C-Terminal Tail of UNC93B1 in Dogs with Exfoliative Cutaneous Lupus Erythematosus (ECLE). <i>Genes</i> , 2020, 11, 159.	1.0	13

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19	Novel protective and risk loci in hip dysplasia in German Shepherds. PLoS Genetics, 2019, 15, e1008197.	1.5	9
20	A comprehensive biomedical variant catalogue based on whole genome sequences of 582 dogs and eight wolves. Animal Genetics, 2019, 50, 695-704.	0.6	138
21	Novel Locus Associated with Symmetrical Lupoid Onychodystrophy in the Bearded Collie. Genes, 2019, 10, 635.	1.0	1
22	Two novel genomic regions associated with fearfulness in dogs overlap human neuropsychiatric loci. Translational Psychiatry, 2019, 9, 18.	2.4	30
23	DLA class II risk haplotypes for autoimmune diseases in the bearded collie offer insight to autoimmunity signatures across dog breeds. Canine Genetics and Epidemiology, 2019, 6, 2.	2.9	15
24	Canine models of human amelogenesis imperfecta: identification of novel recessive ENAM and ACP4 variants. Human Genetics, 2019, 138, 525-533.	1.8	12
25	A frameshift insertion in SGK3 leads to recessive hairlessness in Scottish Deerhounds: a candidate gene for human alopecia conditions. Human Genetics, 2019, 138, 535-539.	1.8	10
26	Genetic dissection of canine hip dysplasia phenotypes and osteoarthritis reveals three novel loci. BMC Genomics, 2019, 20, 1027.	1.2	13
27	A novel <i>KRT71</i> variant in curly-coated dogs. Animal Genetics, 2019, 50, 101-104.	0.6	16
28	A homozygous missense variant in the alkaline phosphatase gene ALPL is associated with a severe form of canine hypophosphatasia. Scientific Reports, 2019, 9, 973.	1.6	11
29	Maternal Inheritance of a Recessive RBP4 Defect in Canine Congenital Eye Disease. Cell Reports, 2018, 23, 2643-2652.	2.9	17
30	Myotonia congenita in a Labrador Retriever with truncated CLCN1. Neuromuscular Disorders, 2018, 28, 597-605.	0.3	12
31	Assembly and Analysis of Unmapped Genome Sequence Reads Reveal Novel Sequence and Variation in Dogs. Scientific Reports, 2018, 8, 10862.	1.6	5
32	MKLN1 splicing defect in dogs with lethal acrodermatitis. PLoS Genetics, 2018, 14, e1007264.	1.5	26
33	Generalized myoclonic epilepsy with photosensitivity in juvenile dogs caused by a defective DIRAS family GTPase 1. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, 2669-2674.	3.3	39
34	Toxoplasma gondii seroprevalence varies by cat breed. PLoS ONE, 2017, 12, e0184659.	1.1	26
35	ANLN truncation causes a familial fatal acute respiratory distress syndrome in Dalmatian dogs. PLoS Genetics, 2017, 13, e1006625.	1.5	14
36	Molecular Characterization of Three Canine Models of Human Rare Bone Diseases: Caffey, van den Ende-Gupta, and Raine Syndromes. PLoS Genetics, 2016, 12, e1006037.	1.5	32

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37	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.	1.1	43
38	Canine models of human rare disorders. Rare Diseases (Austin, Tex), 2016, 4, e1241362.	1.8	37
39	The canine era: the rise of a biomedical model. Animal Genetics, 2016, 47, 519-527.	0.6	44
40	Investigation of rare and low-frequency variants using high-throughput sequencing with pooled DNA samples. Scientific Reports, 2016, 6, 33256.	1.6	13
41	Out of southern East Asia: the natural history of domestic dogs across the world. Cell Research, 2016, 26, 21-33.	5.7	271
42	Canine MPV17 truncation without clinical manifestations. Biology Open, 2015, 4, 1253-1258.	0.6	0
43	Expression of <i>Foxi3</i> is regulated by ectodysplasin in skin appendage placodes. Developmental Dynamics, 2013, 242, 593-603.	0.8	47
44	Ancestral T-Box Mutation Is Present in Many, but Not All, Short-Tailed Dog Breeds. Journal of Heredity, 2009, 100, 236-240.	1.0	34
45	A Mutation in Hairless Dogs Implicates <i>FOXI3</i> in Ectodermal Development. Science, 2008, 321, 1462-1462.	6.0	135
46	BMP-4 affects the differentiation of metanephric mesenchyme and reveals an early anterior-posterior axis of the embryonic kidney. , 2000, 217, 146-158.		120