Ling T Guo

List of Publications by Year
in descending order

26 papers	413 citations	759233 12 h-index	752698 20 g-index
27 all docs	27 docs citations	27 times ranked	533 citing authors

#	Article	IF	Citations
1	Congenital muscular dystrophy in a dog with a <scp>LAMA2</scp> gene deletion. Journal of Veterinary Internal Medicine, 2022, 36, 279-284.	1.6	6
2	Sarcoglycan A mutation in miniature dachshund dogs causes limb-girdle muscular dystrophy 2D. Skeletal Muscle, 2021, 11, 2.	4.2	8
3	Muscular dystrophy-dystroglycanopathy in a family of Labrador retrievers with a LARGE1 mutation. Neuromuscular Disorders, 2021, 31, 1169-1178.	0.6	6
4	Congenital dyserythropoiesis and polymyopathy without cardiac disease in male Labrador retriever littermates. Journal of Veterinary Internal Medicine, 2021, 35, 2409-2414.	1.6	4
5	LAMA2 Nonsense Variant in an Italian Greyhound with Congenital Muscular Dystrophy. Genes, 2021, 12, 1823.	2.4	2
6	Congenital myasthenic syndrome in Golden Retrievers is associated with a novel COLQ mutation. Journal of Veterinary Internal Medicine, 2020, 34, 258-265.	1.6	3
7	A mutation in MTM1 causes X-Linked myotubular myopathy in Boykin spaniels. Neuromuscular Disorders, 2020, 30, 353-359.	0.6	5
8	Pathogenic variants in COL6A3 cause Ullrich-like congenital muscular dystrophy in young Labrador Retriever dogs. Neuromuscular Disorders, 2020, 30, 360-367.	0.6	10
9	Beta-sarcoglycan-deficient muscular dystrophy presenting as chronic bronchopneumonia in a young cat. Journal of Feline Medicine and Surgery Open Reports, 2019, 5, 205511691985645.	0.2	1
10	A Mutation in the Mitochondrial Aspartate/Glutamate Carrier Leads to a More Oxidizing Intramitochondrial Environment and an Inflammatory Myopathy in Dutch Shepherd Dogs. Journal of Neuromuscular Diseases, 2019, 6, 485-501.	2.6	11
11	A glycine transporter SLC6A5 frameshift mutation causes startle disease in Spanish greyhounds. Human Genetics, 2019, 138, 509-513.	3.8	5
12	Myositis, Ganglioneuritis, and Myocarditis with Distinct Perifascicular Muscle Atrophy in a 2-Year-Old Male Boxer. Frontiers in Veterinary Science, 2018, 5, 20.	2.2	0
13	Clinical and genetic characterisation of dystrophin-deficient muscular dystrophy in a family of Miniature Poodle dogs. PLoS ONE, 2018, 13, e0193372.	2.5	16
14	Congenital Myasthenic Syndrome in a Mixed Breed Dog. Frontiers in Veterinary Science, 2017, 4, 173.	2.2	1
15	Exome sequencing reveals independent SGCD deletions causing limb girdle muscular dystrophy in Boston terriers. Skeletal Muscle, 2017, 7, 15.	4.2	18
16	Exome sequencing reveals a nebulin nonsense mutation in a dog model of nemaline myopathy. Mammalian Genome, 2016, 27, 495-502.	2.2	14
17	A CHRNE frameshift mutation causes congenital myasthenic syndrome in young Jack Russell Terriers. Neuromuscular Disorders, 2015, 25, 921-927.	0.6	10
18	X-linked myotubular myopathy in Rottweiler dogs is caused by a missense mutation in Exon 11 of the MTM1 gene. Skeletal Muscle, 2015, 5, 1.	4.2	46

#	Article	IF	Citations
19	Muscular dystrophy in a family of Labrador Retrievers with no muscle dystrophin and a mild phenotype. Neuromuscular Disorders, 2015, 25, 363-370.	0.6	26
20	A COLQ Missense Mutation in Labrador Retrievers Having Congenital Myasthenic Syndrome. PLoS ONE, 2014, 9, e106425.	2.5	17
21	Disruption of both nesprin 1 and desmin results in nuclear anchorage defects and fibrosis in skeletal muscle. Human Molecular Genetics, 2014, 23, 5879-5892.	2.9	52
22	A Novel Mutation in CLCN1 Associated with Feline Myotonia Congenita. PLoS ONE, 2014, 9, e109926.	2.5	17
23	Evaluation of commercial dysferlin antibodies on canine, mouse and human skeletal muscle. Neuromuscular Disorders, 2010, 20, 820-825.	0.6	18
24	Laminin Î±2 (merosin)-deficient muscular dystrophy and demyelinating neuropathy in two cats. Journal of the Neurological Sciences, 2001, 189, 37-43.	0.6	56
25	Muscular Dystrophy in female Dogs. Journal of Veterinary Internal Medicine, 2001, 15, 240-244.	1.6	41
26	Muscular Dystrophy in Female Dogs. Journal of Veterinary Internal Medicine, 2001, 15, 240.	1.6	20