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	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
2	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
3	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
4	Muscular Dystrophy in female Dogs. Journal of Veterinary Internal Medicine, 2001, 15, 240-244.	1.6	41
5	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
6	Muscular Dystrophy in Female Dogs. Journal of Veterinary Internal Medicine, 2001, 15, 240.	1.6	20
7	Evaluation of commercial dysferlin antibodies on canine, mouse and human skeletal muscle. Neuromuscular Disorders, 2010, 20, 820-825.	0.6	18
8	Exome sequencing reveals independent SGCD deletions causing limb girdle muscular dystrophy in Boston terriers. Skeletal Muscle, 2017, 7, 15.	4.2	18
9	A COLQ Missense Mutation in Labrador Retrievers Having Congenital Myasthenic Syndrome. PLoS ONE, 2014, 9, e106425.	2.5	17
10	A Novel Mutation in CLCN1 Associated with Feline Myotonia Congenita. PLoS ONE, 2014, 9, e109926.	2.5	17
11	Clinical and genetic characterisation of dystrophin-deficient muscular dystrophy in a family of Miniature Poodle dogs. PLoS ONE, 2018, 13, e0193372.	2.5	16
12	Exome sequencing reveals a nebulin nonsense mutation in a dog model of nemaline myopathy. Mammalian Genome, 2016, 27, 495-502.	2.2	14
13	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
14	A CHRNE frameshift mutation causes congenital myasthenic syndrome in young Jack Russell Terriers. Neuromuscular Disorders, 2015, 25, 921-927.	0.6	10
15	Pathogenic variants in COL6A3 cause Ullrich-like congenital muscular dystrophy in young Labrador Retriever dogs. Neuromuscular Disorders, 2020, 30, 360-367.	0.6	10
16	Sarcoglycan A mutation in miniature dachshund dogs causes limb-girdle muscular dystrophy 2D. Skeletal Muscle, 2021, 11, 2.	4.2	8
17	Muscular dystrophy-dystroglycanopathy in a family of Labrador retrievers with a LARGE1 mutation. Neuromuscular Disorders, 2021, 31, 1169-1178.	0.6	6
18	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

#	Article	IF	Citations
19	A glycine transporter SLC6A5 frameshift mutation causes startle disease in Spanish greyhounds. Human Genetics, 2019, 138, 509-513.	3.8	5
20	A mutation in MTM1 causes X-Linked myotubular myopathy in Boykin spaniels. Neuromuscular Disorders, 2020, 30, 353-359.	0.6	5
21	Congenital dyserythropoiesis and polymyopathy without cardiac disease in male Labrador retriever littermates. Journal of Veterinary Internal Medicine, 2021, 35, 2409-2414.	1.6	4
22	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
23	LAMA2 Nonsense Variant in an Italian Greyhound with Congenital Muscular Dystrophy. Genes, 2021, 12, 1823.	2.4	2
24	Congenital Myasthenic Syndrome in a Mixed Breed Dog. Frontiers in Veterinary Science, 2017, 4, 173.	2.2	1
25	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
26	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