

Andrew Findlay

List of Publications by Year in descending order

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Version: 2024-02-01

14
papers

527
citations

932766

10
h-index

940134

16
g-index

17
all docs

17
docs citations

17
times ranked

885
citing authors

#	ARTICLE	IF	CITATIONS
1	Clinical utility of anti-cytosolic 5'-nucleotidase 1A antibody in idiopathic inflammatory myopathies. <i>Annals of Clinical and Translational Neurology</i> , 2021, 8, 571-578.	1.7	18
2	Inhibition of DNAJ-HSP70 interaction improves strength in muscular dystrophy. <i>Journal of Clinical Investigation</i> , 2020, 130, 4470-4485.	3.9	14
3	Phenotypic diversity in an international Cure VCP Disease registry. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 267.	1.2	11
4	Lithium chloride corrects weakness and myopathology in a preclinical model of LGMD1D. <i>Neurology: Genetics</i> , 2019, 5, e318.	0.9	15
5	Homozygous recessive MYH2 mutation mimicking dominant MYH2 associated myopathy. <i>Neuromuscular Disorders</i> , 2018, 28, 675-679.	0.3	10
6	Resistance to 6-Methylpurine is Conferred by Defective Adenine Phosphoribosyltransferase in <i>Tetrahymena</i> . <i>Genes</i> , 2018, 9, 179.	1.0	4
7	Efficient Skipping of Single Exon Duplications in DMD Patient-Derived Cell Lines Using an Antisense Oligonucleotide Approach. <i>Journal of Neuromuscular Diseases</i> , 2017, 4, 199-207.	1.1	27
8	The first exon duplication mouse model of Duchenne muscular dystrophy: A tool for therapeutic development. <i>Neuromuscular Disorders</i> , 2015, 25, 827-834.	0.3	27
9	Clinical phenotypes as predictors of the outcome of skipping around <i>DMD</i> exon 45. <i>Annals of Neurology</i> , 2015, 77, 668-674.	2.8	38
10	An overview of polymyositis and dermatomyositis. <i>Muscle and Nerve</i> , 2015, 51, 638-656.	1.0	176
11	Exome sequencing identifies a DNAJB6 mutation in a family with dominantly-inherited limb-girdle muscular dystrophy. <i>Neuromuscular Disorders</i> , 2014, 24, 431-435.	0.3	35
12	The ZZ Domain of Dystrophin in DMD: Making Sense of Missense Mutations. <i>Human Mutation</i> , 2014, 35, 257-264.	1.1	23
13	Translation from a DMD exon 5 IRES results in a functional dystrophin isoform that attenuates dystrophinopathy in humans and mice. <i>Nature Medicine</i> , 2014, 20, 992-1000.	15.2	113
14	Camptocormia as a late presentation in a manifesting carrier of duchenne muscular dystrophy. <i>Muscle and Nerve</i> , 2013, 47, 124-127.	1.0	8