Andrew Findlay

List of Publications by Year
in descending order

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