Nicolas Wein

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/11700880/publications.pdf

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25	745	567281	610901
papers	citations	15	24
		h-index	g-index

28	28	28	898
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Intron mutations and early transcription termination in Duchenne and Becker muscular dystrophy. Human Mutation, 2022, 43, 511-528.	2.5	16
2	Designed U7 snRNAs inhibitÂDUX4Âexpression and improve FSHD-associated outcomes inÂDUX4Âoverexpressing cells and FSHD patient myotubes. Molecular Therapy - Nucleic Acids, 2021, 23, 476-486.	5.1	17
3	Direct Reprogramming of Human Fibroblasts into Myoblasts to Investigate Therapies for Neuromuscular Disorders. Journal of Visualized Experiments, 2021, , .	0.3	0
4	Pre-clinical dose-escalation studies establish a therapeutic range for U7snRNA-mediated DMD exon 2 skipping. Molecular Therapy - Methods and Clinical Development, 2021, 21, 325-340.	4.1	21
5	Absence of Significant Off-Target Splicing Variation with a U7snRNA Vector Targeting DMD Exon 2 Duplications. Human Gene Therapy, 2021, 32, 1346-1359.	2.7	8
6	U7 snRNA, a Small RNA with a Big Impact in Gene Therapy. Human Gene Therapy, 2021, 32, 1317-1329.	2.7	5
7	Lack of Toxicity in Nonhuman Primates Receiving Clinically Relevant Doses of an AAV9.U7snRNA Vector Designed to Induce DMD Exon 2 Skipping. Human Gene Therapy, 2021, 32, 882-894.	2.7	29
8	X-linked muscular dystrophy in a Labrador Retriever strain: phenotypic and molecular characterisation. Skeletal Muscle, 2020, 10, 23.	4.2	12
9	Alternate Translational Initiation of Dystrophin: A Novel Therapeutic Approach. , 2019, , 371-382.		0
10	Personalized gene and cell therapy for Duchenne Muscular Dystrophy. Neuromuscular Disorders, 2018, 28, 803-824.	0.6	45
11	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.	2.6	27
12	Dp412e: a novel human embryonic dystrophin isoform induced by BMP4 in early differentiated cells. Skeletal Muscle, 2015, 5, 40.	4.2	28
13	Exon 32 Skipping of Dysferlin Rescues Membrane Repair in Patientsâ€™ Cells. Journal of Neuromuscular Diseases, 2015, 2, 281-290.	2.6	29
14	The first exon duplication mouse model of Duchenne muscular dystrophy: A tool for therapeutic development. Neuromuscular Disorders, 2015, 25, 827-834.	0.6	27
15	Clinical phenotypes as predictors of the outcome of skipping around <scp>DMD</scp> exon 45. Annals of Neurology, 2015, 77, 668-674.	5.3	38
16	Genetics and Emerging Treatments for Duchenne and Becker Muscular Dystrophy. Pediatric Clinics of North America, 2015, 62, 723-742.	1.8	71
17	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.	30.7	113
18	Becker muscular dystrophy with widespread muscle hypertrophy and a non-sense mutation of exon 2. Neuromuscular Disorders, 2013, 23, 192.	0.6	1

#	Article	IF	Citations
19	Fullâ€length dysferlin expression driven by engineered human dystrophic blood derived <scp>CD</scp>133+ stem cells. FEBS Journal, 2013, 280, 6045-6060.	4.7	12
20	UMD-DYSF, a novel locus specific database for the compilation and interactive analysis of mutations in the dysferlin gene. Human Mutation, 2012, 33, E2317-E2331.	2.5	35
21	Translational Research and Therapeutic Perspectives in Dysferlinopathies. Molecular Medicine, 2011, 17, 875-882.	4.4	36
22	Efficient bypass of mutations in dysferlin deficient patient cells by antisense-induced exon skipping. Human Mutation, 2010, 31, 136-142.	2.5	80
23	Therapeutic exon â€˜switchingâ€™ for dysferlinopathies?. European Journal of Human Genetics, 2010, 18, 969-970.	2.8	7
24	A Naturally Occurring Human Minidysferlin Protein Repairs Sarcolemmal Lesions in a Mouse Model of Dysferlinopathy. Science Translational Medicine, 2010, 2, 50ra69.	12.4	69
25	Identification of Different Genomic Deletions and One Duplication in the Dysferlin Gene Using Multiplex Ligation-Dependent Probe Amplification and Genomic Quantitative PCR. Genetic Testing and Molecular Biomarkers, 2009, 13, 439-442.	0.7	18