

Daniel G Miller

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5124360/publications.pdf>

Version: 2024-02-01

44
papers

6,739
citations

147801

31
h-index

254184

43
g-index

45
all docs

45
docs citations

45
times ranked

5594
citing authors

#	ARTICLE	IF	CITATIONS
1	A Unifying Genetic Model for Facioscapulohumeral Muscular Dystrophy. <i>Science</i> , 2010, 329, 1650-1653.	12.6	638
2	Cell-surface receptors for gibbon ape leukemia virus and amphotropic murine retrovirus are inducible sodium-dependent phosphate symporters.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1994, 91, 7071-7075.	7.1	623
3	Systemic delivery of genes to striated muscles using adeno-associated viral vectors. <i>Nature Medicine</i> , 2004, 10, 828-834.	30.7	586
4	Digenic inheritance of an SMCHD1 mutation and an FSHD-permissive D4Z4 allele causes facioscapulohumeral muscular dystrophy type 2. <i>Nature Genetics</i> , 2012, 44, 1370-1374.	21.4	582
5	AAV Vector Integration Sites in Mouse Hepatocellular Carcinoma. <i>Science</i> , 2007, 317, 477-477.	12.6	532
6	Cloning of the cellular receptor for amphotropic murine retroviruses reveals homology to that for gibbon ape leukemia virus.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1994, 91, 78-82.	7.1	407
7	[40] Use of retroviral vectors for gene transfer and expression. <i>Methods in Enzymology</i> , 1993, 217, 581-599.	1.0	395
8	Facioscapulohumeral Dystrophy: Incomplete Suppression of a Retrotransposed Gene. <i>PLoS Genetics</i> , 2010, 6, e1001181.	3.5	394
9	Adeno-associated virus vectors integrate at chromosome breakage sites. <i>Nature Genetics</i> , 2004, 36, 767-773.	21.4	226
10	Foamy virus vector integration sites in normal human cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2006, 103, 1498-1503.	7.1	226
11	RNA transcripts, miRNA-sized fragments and proteins produced from D4Z4 units: new candidates for the pathophysiology of facioscapulohumeral dystrophy. <i>Human Molecular Genetics</i> , 2009, 18, 2414-2430.	2.9	182
12	Endogenous DUX4 expression in FSHD myotubes is sufficient to cause cell death and disrupts RNA splicing and cell migration pathways. <i>Human Molecular Genetics</i> , 2015, 24, 5901-5914.	2.9	159
13	Characterization of microRNAs Involved in Embryonic Stem Cell States. <i>Stem Cells and Development</i> , 2010, 19, 935-950.	2.1	156
14	Chromosomal effects of adeno-associated virus vector integration. <i>Nature Genetics</i> , 2002, 30, 147-148.	21.4	148
15	Large-Scale Analysis of Adeno-Associated Virus Vector Integration Sites in Normal Human Cells. <i>Journal of Virology</i> , 2005, 79, 11434-11442.	3.4	148
16	Inter-individual differences in CpG methylation at D4Z4 correlate with clinical variability in FSHD1 and FSHD2. <i>Human Molecular Genetics</i> , 2015, 24, 659-669.	2.9	130
17	Human Gene Targeting by Adeno-Associated Virus Vectors Is Enhanced by DNA Double-Strand Breaks. <i>Molecular and Cellular Biology</i> , 2003, 23, 3550-3557.	2.3	123
18	Comparison of HIV-derived Lentiviral and MLV-based Gammaretroviral Vector Integration Sites in Primate Repopulating Cells. <i>Molecular Therapy</i> , 2007, 15, 1356-1365.	8.2	104

#	ARTICLE	IF	CITATIONS
19	Gene targeting in vivo by adeno-associated virus vectors. <i>Nature Biotechnology</i> , 2006, 24, 1022-1026.	17.5	102
20	A Human Pluripotent Stem Cell Model of Facioscapulohumeral Muscular Dystrophy-Affected Skeletal Muscles. <i>Stem Cells Translational Medicine</i> , 2016, 5, 1145-1161.	3.3	98
21	Wnt/ β -catenin signaling suppresses DUX4 expression and prevents apoptosis of FSHD muscle cells. <i>Human Molecular Genetics</i> , 2013, 22, 4661-4672.	2.9	92
22	GCC Repeat Expansion and Exon 1 Methylation of XYL1 Is a Common Pathogenic Variant in Baratela-Scott Syndrome. <i>American Journal of Human Genetics</i> , 2019, 104, 35-44.	6.2	81
23	Unique Integration Profiles in a Canine Model of Long-Term Repopulating Cells Transduced with Gammaretrovirus, Lentivirus, or Foamy Virus. <i>Human Gene Therapy</i> , 2007, 18, 423-434.	2.7	73
24	Frequent Endonuclease Cleavage at Off-target Locations In Vivo. <i>Molecular Therapy</i> , 2010, 18, 983-986.	8.2	54
25	Longitudinal features of stir bright signal in FSHD¹. <i>Muscle and Nerve</i> , 2014, 49, 257-260.	2.2	46
26	Efficient KRT14 Targeting and Functional Characterization of Transplanted Human Keratinocytes for the Treatment of Epidermolysis Bullosa Simplex. <i>Molecular Therapy</i> , 2010, 18, 1624-1632.	8.2	43
27	Clinical trial preparedness in facioscapulohumeral muscular dystrophy: Clinical, tissue, and imaging outcome measures 29â€“30 May 2015, Rochester, New York. <i>Neuromuscular Disorders</i> , 2016, 26, 181-186.	0.6	43
28	Facioscapulohumeral muscular dystrophy. <i>Current Opinion in Neurology</i> , 2012, 25, 614-620.	3.6	42
29	Chromatin structure of two genomic sites for targeted transgene integration in induced pluripotent stem cells and hematopoietic stem cells. <i>Gene Therapy</i> , 2013, 20, 201-214.	4.5	39
30	VISA - Vector Integration Site Analysis server: a web-based server to rapidly identify retroviral integration sites from next-generation sequencing. <i>BMC Bioinformatics</i> , 2015, 16, 212.	2.6	37
31	A cross sectional study of two independent cohorts identifies serum biomarkers for facioscapulohumeral muscular dystrophy (FSHD). <i>Neuromuscular Disorders</i> , 2016, 26, 405-413.	0.6	36
32	MRI change metrics of facioscapulohumeral muscular dystrophy: Stir and T1. <i>Muscle and Nerve</i> , 2018, 57, 905-912.	2.2	29
33	Sporadic DUX4 expression in FSHD myocytes is associated with incomplete repression by the PRC2 complex and gain of H3K9 acetylation on the contracted D4Z4 allele. <i>Epigenetics and Chromatin</i> , 2018, 11, 47.	3.9	26
34	Gene Therapy for Hemophilia. <i>New England Journal of Medicine</i> , 2001, 344, 1782-1784.	27.0	22
35	Asymmetric Bidirectional Transcription from the FSHD-Causing D4Z4 Array Modulates DUX4 Production. <i>PLoS ONE</i> , 2012, 7, e35532.	2.5	20
36	Expression patterns of FSHD-causing DUX4 and myogenic transcription factors PAX3 and PAX7 are spatially distinct in differentiating human stem cell cultures. <i>Skeletal Muscle</i> , 2017, 7, 13.	4.2	17

#	ARTICLE	IF	CITATIONS
37	Epigenetic memory via concordant DNA methylation is inversely correlated to developmental potential of mammalian cells. PLoS Genetics, 2017, 13, e1007060.	3.5	17
38	Efficient PRNP Gene Targeting in Bovine Fibroblasts by Adeno-Associated Virus Vectors. Cloning and Stem Cells, 2004, 6, 31-36.	2.6	16
39	Epigenetic regulation of the X-chromosomal macrosatellite repeat encoding for the cancer/testis gene CT47. European Journal of Human Genetics, 2012, 20, 185-191.	2.8	15
40	Integration Bias of Gamma-retrovirus Vectors following Transduction and Growth of Primary Mouse Hematopoietic Progenitor Cells with and without Selection. Molecular Therapy, 2006, 14, 226-235.	8.2	14
41	AAV-Mediated Gene Targeting. Methods in Molecular Biology, 2012, 807, 301-315.	0.9	9
42	Small noncoding RNAs in FSHD2 muscle cells reveal both DUX4- and SMCHD1-specific signatures. Human Molecular Genetics, 2018, 27, 2644-2657.	2.9	6
43	Underperforming Big Ideas in Biomedical Research. JAMA - Journal of the American Medical Association, 2017, 317, 321.	7.4	2
44	Gene Therapy for Facioscapulohumeral Muscular Dystrophy (FSHD). , 2019, , 509-524.		1