

Dan Wang

List of Publications by Citations

Source: <https://exaly.com/author-pdf/1954452/dan-wang-publications-by-citations.pdf>

Version: 2024-04-27

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

37
papers

1,873
citations

21
h-index

43
g-index

52
ext. papers

2,620
ext. citations

12.4
avg, IF

5.66
L-index

#	Paper	IF	Citations
37	Adeno-associated virus vector as a platform for gene therapy delivery. <i>Nature Reviews Drug Discovery</i> , 2019 , 18, 358-378	64.1	555
36	Adenovirus-Mediated Somatic Genome Editing of Pten by CRISPR/Cas9 in Mouse Liver in Spite of Cas9-Specific Immune Responses. <i>Human Gene Therapy</i> , 2015 , 26, 432-42	4.8	226
35	CRISPR-Based Therapeutic Genome Editing: Strategies and In Vivo Delivery by AAV Vectors. <i>Cell</i> , 2020 , 181, 136-150	56.2	137
34	AMH/MIS as a contraceptive that protects the ovarian reserve during chemotherapy. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017 , 114, E1688-E1697	11.5	98
33	Attenuation of nonsense-mediated mRNA decay enhances in vivo nonsense suppression. <i>PLoS ONE</i> , 2013 , 8, e60478	3.7	77
32	Suppression of premature termination codons as a therapeutic approach. <i>Critical Reviews in Biochemistry and Molecular Biology</i> , 2012 , 47, 444-63	8.7	72
31	The potential of adeno-associated viral vectors for gene delivery to muscle tissue. <i>Expert Opinion on Drug Delivery</i> , 2014 , 11, 345-364	8	65
30	The designer aminoglycoside NB84 significantly reduces glycosaminoglycan accumulation associated with MPS I-H in the Idua-W392X mouse. <i>Molecular Genetics and Metabolism</i> , 2012 , 105, 116-23	3.7	62
29	State-of-the-art human gene therapy: part II. Gene therapy strategies and clinical applications. <i>Discovery Medicine</i> , 2014 , 18, 151-61	2.5	57
28	State-of-the-art human gene therapy: part I. Gene delivery technologies. <i>Discovery Medicine</i> , 2014 , 18, 67-77	2.5	50
27	CNS-restricted Transduction and CRISPR/Cas9-mediated Gene Deletion with an Engineered AAV Vector. <i>Molecular Therapy - Nucleic Acids</i> , 2016 , 5, e338	10.7	48
26	Streamlined ex vivo and in vivo genome editing in mouse embryos using recombinant adeno-associated viruses. <i>Nature Communications</i> , 2018 , 9, 412	17.4	41
25	In Vivo Genome Editing Partially Restores Alpha1-Antitrypsin in a Murine Model of AAT Deficiency. <i>Human Gene Therapy</i> , 2018 , 29, 853-860	4.8	40
24	Short DNA Hairpins Compromise Recombinant Adeno-Associated Virus Genome Homogeneity. <i>Molecular Therapy</i> , 2017 , 25, 1363-1374	11.7	36
23	AAV9 delivering a modified human Mullerian inhibiting substance as a gene therapy in patient-derived xenografts of ovarian cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015 , 112, E4418-27	11.5	34
22	Characterization of an MPS I-H knock-in mouse that carries a nonsense mutation analogous to the human IDUA-W402X mutation. <i>Molecular Genetics and Metabolism</i> , 2010 , 99, 62-71	3.7	33
21	Bone-targeting AAV-mediated silencing of Schnurri-3 prevents bone loss in osteoporosis. <i>Nature Communications</i> , 2019 , 10, 2958	17.4	29

20	Adeno-Associated Virus Neutralizing Antibodies in Large Animals and Their Impact on Brain Intraparenchymal Gene Transfer. <i>Molecular Therapy - Methods and Clinical Development</i> , 2018 , 11, 65-72	6.4	27
19	A Rationally Engineered Capsid Variant of AAV9 for Systemic CNS-Directed and Peripheral Tissue-Detargeted Gene Delivery in Neonates. <i>Molecular Therapy - Methods and Clinical Development</i> , 2018 , 9, 234-246	6.4	26
18	Cas9-mediated allelic exchange repairs compound heterozygous recessive mutations in mice. <i>Nature Biotechnology</i> , 2018 , 36, 839-842	44.5	23
17	A Single Injection of Recombinant Adeno-Associated Virus into the Lumbar Cistern Delivers Transgene Expression Throughout the Whole Spinal Cord. <i>Molecular Neurobiology</i> , 2016 , 53, 3235-3248	6.2	22
16	Single-cell sequencing of neonatal uterus reveals an Msr2+ endometrial progenitor indispensable for fertility. <i>ELife</i> , 2019 , 8,	8.9	18
15	Slow Intrathecal Injection of rAAVrh10 Enhances its Transduction of Spinal Cord and Therapeutic Efficacy in a Mutant SOD1 Model of ALS. <i>Neuroscience</i> , 2017 , 365, 192-205	3.9	14
14	Circumventing cellular immunity by miR142-mediated regulation sufficiently supports rAAV-delivered OVA expression without activating humoral immunity. <i>JCI Insight</i> , 2019 , 5,	9.9	14
13	AAV-Genome Population Sequencing of Vectors Packaging CRISPR Components Reveals Design-Influenced Heterogeneity. <i>Molecular Therapy - Methods and Clinical Development</i> , 2020 , 18, 639-651	6.4	12
12	Intrathecal Adeno-Associated Viral Vector-Mediated Gene Delivery for Adrenomyeloneuropathy. <i>Human Gene Therapy</i> , 2019 , 30, 544-555	4.8	12
11	Bone-Targeting AAV-Mediated Gene Silencing in Osteoclasts for Osteoporosis Therapy. <i>Molecular Therapy - Methods and Clinical Development</i> , 2020 , 17, 922-935	6.4	8
10	Adeno-associated Virus Serotype Vectors Efficiently Transduce Normal Prostate Tissue and Prostate Cancer Cells. <i>European Urology</i> , 2016 , 69, 179-81	10.2	7
9	Self-inactivating, all-in-one AAV vectors for precision Cas9 genome editing via homology-directed repair in vivo. <i>Nature Communications</i> , 2021 , 12, 6267	17.4	5
8	Single-cell sequencing reveals suppressive transcriptional programs regulated by MIS/AMH in neonatal ovaries. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021 , 118,	11.5	5
7	Gene Delivery to Nonhuman Primate Preimplantation Embryos Using Recombinant Adeno-Associated Virus. <i>Advanced Science</i> , 2019 , 6, 1900440	13.6	4
6	Slow Infusion of Recombinant Adeno-Associated Viruses into the Mouse Cerebrospinal Fluid Space. <i>Human Gene Therapy Methods</i> , 2018 , 29, 75-85	4.9	3
5	AAV-delivered suppressor tRNA overcomes a nonsense mutation in mice.. <i>Nature</i> , 2022 ,	50.4	3
4	Taking a Hint from Structural Biology: To Better Understand AAV Transport across the BBB. <i>Molecular Therapy</i> , 2018 , 26, 336-338	11.7	2
3	AAV-Mediated Gene Therapy for Glycosphingolipid Biosynthesis Deficiencies. <i>Trends in Molecular Medicine</i> , 2021 , 27, 520-523	11.5	0

2 Viral Vectors for Muscle Gene Therapy **2019**, 179-192

1 Pharmacological suppression of nonsense mutations to treat genetic diseases. *FASEB Journal*, **2010**, 24, 187.2

0.9