

# Alan McClelland

## List of Publications by Year in descending order

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Version: 2024-02-01

28  
papers

5,173  
citations

448610

19  
h-index

591227

27  
g-index

28  
all docs

28  
docs citations

28  
times ranked

3193  
citing authors

#	ARTICLE	IF	CITATIONS
1	The major human rhinovirus receptor is ICAM-1. <i>Cell</i> , 1989, 56, 839-847.	13.5	1,170
2	Evidence for gene transfer and expression of factor IX in haemophilia B patients treated with an AAV vector. <i>Nature Genetics</i> , 2000, 24, 257-261.	9.4	971
3	AAV-mediated factor IX gene transfer to skeletal muscle in patients with severe hemophilia B. <i>Blood</i> , 2003, 101, 2963-2972.	0.6	707
4	Adenovirus mediated expression of therapeutic plasma levels of human factor IX in mice. <i>Nature Genetics</i> , 1993, 5, 397-402.	9.4	376
5	The human transferrin receptor gene: genomic organization, and the complete primary structure of the receptor deduced from a cDNA sequence. <i>Cell</i> , 1984, 39, 267-274.	13.5	360
6	Gene transfer, expression, and molecular cloning of the human transferrin receptor gene. <i>Cell</i> , 1984, 37, 95-103.	13.5	236
7	Preclinical in vivo evaluation of pseudotyped adeno-associated virus vectors for liver gene therapy. <i>Blood</i> , 2003, 102, 2412-2419.	0.6	196
8	Quantification of adeno-associated virus particles and empty capsids by optical density measurement. <i>Molecular Therapy</i> , 2003, 7, 122-128.	3.7	172
9	Expression of Tissue Inhibitor of Matrix Metalloproteinases 1 by Use of an Adenoviral Vector Inhibits Smooth Muscle Cell Migration and Reduces Neointimal Hyperplasia in the Rat Model of Vascular Balloon Injury. <i>Circulation</i> , 1999, 99, 3199-3205.	1.6	118
10	Sustained Phenotypic Correction of Murine Hemophilia A by In Vivo Gene Therapy. <i>Blood</i> , 1998, 91, 3273-3281.	0.6	111
11	Sustained phenotypic correction of canine hemophilia A using an adeno-associated viral vector. <i>Blood</i> , 2003, 102, 2031-2037.	0.6	101
12	Circumvention of Immunity to the Adenovirus Major Coat Protein Hexon. <i>Journal of Virology</i> , 1998, 72, 6875-6879.	1.5	99
13	Striatal Delivery of rAAV-hAADC to Rats with Preexisting Immunity to AAV. <i>Molecular Therapy</i> , 2004, 9, 403-409.	3.7	89
14	<i>In Vivo</i> Gene Delivery and Expression of Physiological Levels of Functional Human Factor VIII in Mice. <i>Human Gene Therapy</i> , 1995, 6, 185-193.	1.4	87
15	High-Level Tissue-Specific Expression of Functional Human Factor VIII in Mice. <i>Human Gene Therapy</i> , 1996, 7, 183-195.	1.4	77
16	The molecular cloning of a dispersed set of developmentally regulated genes which encode the major larval serum protein of <i>D. melanogaster</i> . <i>Cell</i> , 1981, 23, 441-449.	13.5	71
17	Phenotypic correction of a mouse model of hemophilia A using AAV2 vectors encoding the heavy and light chains of FVIII. <i>Blood</i> , 2003, 102, 3919-3926.	0.6	67
18	Phenotypic Correction of Hypercholesterolemia in ApoE-Deficient Mice by Adenovirus-Mediated In Vivo Gene Transfer. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 1995, 15, 479-484.	1.1	56

#	ARTICLE	IF	CITATIONS
19	Sequence conservation around the 5' ends of the larval serum protein 1 genes of <i>Drosophila melanogaster</i> . <i>Journal of Molecular Biology</i> , 1986, 189, 1-11.	2.0	42
20	Preliminary X-ray crystallographic analysis of intercellular adhesion molecule-1. <i>Journal of Molecular Biology</i> , 1992, 225, 1127-1130.	2.0	20
21	Short intervening sequences close to the 5' ends of the three <i>Drosophila</i> larval serum protein 1 genes. <i>Journal of Molecular Biology</i> , 1981, 153, 257-272.	2.0	16
22	Enhanced Neuroblastoma Transduction for an Improved Antitumor Vaccine. <i>Journal of Surgical Research</i> , 1999, 83, 95-99.	0.8	8
23	In Vivo Adenoviral Gene Transfer of TIMP-1 after Vascular Injury Reduces Neointimal Formation. <i>Annals of the New York Academy of Sciences</i> , 1999, 878, 742-743.	1.8	7
24	Sustained Phenotypic Correction of Murine Hemophilia A by In Vivo Gene Therapy. <i>Blood</i> , 1998, 91, 3273-3281.	0.6	7
25	[25] Molecular cloning of receptor genes by transfection. <i>Methods in Enzymology</i> , 1987, 147, 280-291.	0.4	6
26	Linkage of a <i>Drosophila melanogaster</i> U1 small nuclear RNA gene to the larval serum protein 1 gene. <i>Journal of Molecular Biology</i> , 1985, 185, 649.	2.0	1
27	Adenoviral Gene Delivery Approaches for Systemic Expression. <i>Developments in Cardiovascular Medicine</i> , 1997, , 433-448.	0.1	1
28	Structure of a Human Rhinovirus Complexed with its Receptor Molecule. , 1993, , 1-12.		1