

# Successful transduction of liver in hemophilia by AAV- $\mu$ the host immune response

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Gene therapy for viral hepatitis. <i>Expert Opinion on Biological Therapy</i> , 2006, 6, 1263-1278.	1.4	5
2	Long-Term Increase in mVEGF164 in Mouse Hindlimb Muscle Mediated by Phage $\lambda$ C31 Integrase After Nonviral DNA Delivery. <i>Human Gene Therapy</i> , 2006, 17, 871-876.	1.4	32
3	Sleeping Beauty Transposon-Mediated Nonviral Gene Therapy. <i>BioDrugs</i> , 2006, 20, 219-229.	2.2	14
4	Gene therapy for treatment of inherited haematological disorders. <i>Expert Opinion on Biological Therapy</i> , 2006, 6, 509-522.	1.4	13
5	Long-Term Efficacy of Adeno-Associated Virus Serotypes 8 and 9 in Hemophilia A Dogs and Mice. <i>Human Gene Therapy</i> , 2006, 17, 427-439.	1.4	95
6	AAV-mediated gene transfer for retinal diseases. <i>Expert Opinion on Biological Therapy</i> , 2006, 6, 1279-1294.	1.4	36
8	Evidence of Multiyear Factor IX Expression by AAV-Mediated Gene Transfer to Skeletal Muscle in an Individual with Severe Hemophilia B. <i>Molecular Therapy</i> , 2006, 14, 452-455.	3.7	196
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21	Genetic Modification of Somatic Gut Mucosa. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2006, 43, 158-159.	0.9	2

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1278	Development of AAV Variants with Human Hepatocyte Tropism and Neutralizing Antibody Escape Capacity. <i>Molecular Therapy - Methods and Clinical Development</i> , 2020, 18, 259-268.	1.8	20
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1324	Capsid-specific removal of circulating antibodies to adeno-associated virus vectors. <i>Scientific Reports</i> , 2020, 10, 864.	1.6	72
1326	Muscle-Directed Delivery of an AAV1 Vector Leads to Capsid-Specific T Cell Exhaustion in Nonhuman Primates and Humans. <i>Molecular Therapy</i> , 2020, 28, 747-757.	3.7	23
1327	Translational Potential of Immune Tolerance Induction by AAV Liver-Directed Factor VIII Gene Therapy for Hemophilia A. <i>Frontiers in Immunology</i> , 2020, 11, 618.	2.2	22
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