

Successful transduction of liver in hemophilia by AAV-P_{EF} the host immune response

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Gene therapy for viral hepatitis. Expert Opinion on Biological Therapy, 2006, 6, 1263-1278.	3.1	5
2	Long-Term Increase in mVEGF164 in Mouse Hindlimb Muscle Mediated by Phage- λ C31 Integrase After Nonviral DNA Delivery. Human Gene Therapy, 2006, 17, 871-876.	2.7	32
3	Sleeping Beauty Transposon-Mediated Nonviral Gene Therapy. BioDrugs, 2006, 20, 219-229.	4.6	14
4	Gene therapy for treatment of inherited haematological disorders. Expert Opinion on Biological Therapy, 2006, 6, 509-522.	3.1	13
5	Long-Term Efficacy of Adeno-Associated Virus Serotypes 8 and 9 in Hemophilia A Dogs and Mice. Human Gene Therapy, 2006, 17, 427-439.	2.7	95
6	AAV-mediated gene transfer for retinal diseases. Expert Opinion on Biological Therapy, 2006, 6, 1279-1294.	3.1	36
8	Evidence of Multiyear Factor IX Expression by AAV-Mediated Gene Transfer to Skeletal Muscle in an Individual with Severe Hemophilia B. Molecular Therapy, 2006, 14, 452-455.	8.2	196
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