

nAAVigen Therapeutics

Scientific Publication Directory

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Hematopoietic Stem/Progenitor Cell Transduction with AAV Vectors

1. A. Srivastava. Parvovirus-based vectors for human gene therapy. *Blood Cells*, 20: 531- 538, 1994.
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Human Globin Gene Transfer with AAV Vectors

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AAV-B19 Hybrid and Chimeric Vectors

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Decoy Zinc Finger-Binding Proteins

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Supporting Publications

1. K.Y. Qing, X.-S. Wang, D.M. Kube, S. Ponnazhagan, A. Bajpai and A. Srivastava. Role of tyrosine phosphorylation of a cellular protein in adeno-associated virus 2-mediated transgene expression. *Proc. Natl. Acad. Sci., USA*, 94: 10879-10884, 1997.
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21. B. Li, W. Ma, G.V. Aslanidi, C. Ling, K.V. Vliet, L.-y. Huang, M. Agbandje-McKenna, A. Srivastava, and G.A. Aslanidi. Site-directed mutagenesis of surface-exposed lysine residues leads to high-efficiency transduction by recombinant AAV2, but not AAV8 vectors. *Hum. Gene Therapy Methods*, 26: 211-220, 2015.