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Gene therapy is the (sometimes called biological nanoparticles or viral vectors) and naked DNA the therapeutic DNA introduced into target cells must

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The use of synthetic oligonucleotides in gene therapy is to The oligonucleotide is designed to anneal with complementarity to the target gene with

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Viral vectors: from virology to transgene

Keywords: viral vectors, gene therapy, AAV, Ad, HSV, lentivectors, pseudotyping, cell targeting, transcriptional targeting, vectors production

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