

Enhanced Safety High Titers

AAVPrime™ Adeno-associated Viral (AAV) Particles

For *in vitr*o and *in vivo* studies

Advantages

- Titer of purified AAV particles can be up to 10¹⁴ GC/ml
- All serotypes available allows tissue selectivity
- Purified particles for in vivo animal studies
- Low toxicity and minimal host immune response
- Custom packaging service for ORF, shRNA, CRISPR and more

Primary Target Tissues									
Serotype	Retina	Neurons	Brain	Lung	Heart	Liver	Muscle	Kidney	Pancreas
AAV-1		\checkmark			\checkmark		\checkmark		\checkmark
AAV-2	\checkmark	\checkmark	\checkmark			\checkmark	\checkmark	\checkmark	
AAV-3	\checkmark			\checkmark		\checkmark	\checkmark		
AAV-4	\checkmark	\checkmark	\checkmark				\checkmark		
AAV-5	\checkmark	\checkmark		\checkmark					
AAV-6				\checkmark	\checkmark	\checkmark	\checkmark		
AAV-7	\checkmark	\checkmark				\checkmark	\checkmark		\checkmark
AAV-8	\checkmark		\checkmark			\checkmark	\checkmark		
AAV-9			\checkmark						
AAV-10		\checkmark		\checkmark	\checkmark	\checkmark	\checkmark		
AAV-DJ	Efficiently transduces a wide variety of cell types in vitro								
AAV-DJ/8	A variant of AAV-DJ that permits infection of liver as well as other tissues in vivo								

AAV Serotype Testing Kit

GeneCopoeia's AAVPrime[™] Adeno-associated virus (AAV) Serotype Testing Kit contains 9 premade GFP-expressing AAV in serotypes 1, 2, 3, 6, 7, 8, 9, DJ and DJ/8. The kit can be used to determine the ideal AAV serotype for the infection of different tissue or cell types.



AAV of different serotypes carrying GFP were used to transduce HT1080 cell line at MOI 20,000 as shown in the figure above. Fluorescent images (GFP) show that cells are transduced with AAV-GFP vectors from the AAV Serotype Testing Kit.

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Applications: For *in vivo* and *in vitro* studies of gene overexpression, knockdown and knockout. Type of custom AAV service available: ORF cDNA, shRNA, CRISPR sgRNA and SaCas9. Control AAV particles such as eGFP, mCherry, LacZ, luciferase, etc. in various serotypes are also available.



Figure 1. HT1080 cells in 24-well plates were transduced with 0.5 μ L of standard AAV particles (MOI=200) expressing either eGFP (A), RFP (B), or mCherry (C). Cells were visualized with a fluorescence microscope 48 hours post-transduction (Exposure time: 400 ms).



Figure 2. HT1080 cells in 24-well plates were transduced with the indicated amounts of purified AAV particles (MOI=100, 1000, 10000) expressing eGFP. Cells were visualized with a fluorescence microscope 72 hours post-transduction (Exposure time: 400 ms).

Figure 3. HT1080 cells in 24-well plates were co-transduced standard AAV particles AAV-2-eGFP expressing eGFP and AAV-2-shRNA knocking down eGFP with series of MOI ratios, 3000:0(A); 3000:1500(B); 3000:3000(C) and 3000:6000(D). Cells were visualized with a fluorescence microscope 48 hours post-transduction (Exposure time: 400 ms).

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