

Product datasheet

anti-AAV9 (intact particle) mouse monoclonal, ADK9, liquid, purified, sample

Short overview

Cat. No.	690162S
Quantity	200 µl (50 µg/ml)
Concentration	50 µg/ml

Product description

Host	Mouse
Antibody Type	Monoclonal
Isotype	IgA kappa
Conjugate	Unconjugated
Purification	Affinity chromatography
Storage	Up to 1 month: 2-8°C; long term storage in aliquots at -20°C; avoid freeze/thaw cycles
Intended use	Research use only
Application	Dot blot, ELISA, ICC/IF, Neutralization assay
Reactivity	AAV9
No reactivity	AAV1, AAV11, AAV12, AAV2, AAV3, AAV4, AAV5, AAV6, AAV7, AAV8, AAVDJ, AAVrh10, AAVrh74

Applications

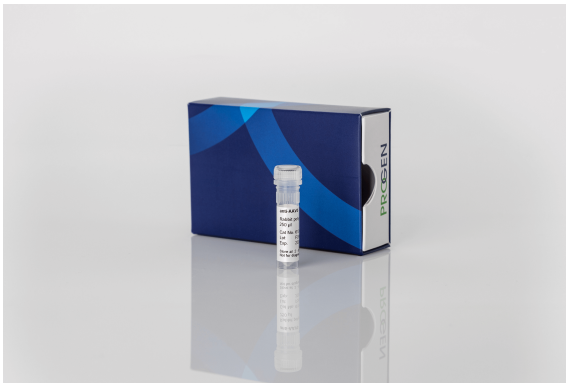
Dot Blot	1:100-1:1,000 (0.05-0.5 µg/ml; non-denaturing conditions)
ELISA	Assay dependent
Immunocytochemistry (ICC)	Assay dependent
Neutralization Assay	EC50 ~2 ng/ml (AAV9) - assay dependent

Background

For characterization of different stages of infection and very useful for the analysis of the AAV assembly process. ADK9 specifically reacts with intact adeno-associated virus particles, empty and full capsids. Recognizes a conformational epitope of assembled capsids. The antibody cannot be used for immunoblotting. The antibody is also useful for neutralizing experiments.

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Product images



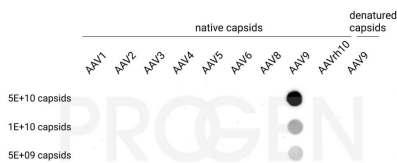
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Serotype	Clone	Residues	
AAV9	ADK9	453-GSGQN-457	Neutralization epitope, not sure because no other epitope was i

Adachi, K., Enoki, T., Kawano, Y., Veraz, M. & Nakai, H. Drawing a high-resolution functional map of adeno-sequencing. *Nat. Commun.* 5, (2014).

For AAV9 the neutralization epitope has been identified and might give an additional indication about the binding residues of the neutralizing antibody ADK9 used for the corresponding AAV9 Titration ELISAs. ADK9 recognizes and neutralizes assembled AAV9 capsids.

Adachi, K., Enoki, T., Kawano, Y., Veraz, M. & Nakai, H. Drawing a high-resolution functional map of adeno-associated virus capsid by massively parallel sequencing. *Nat. Commun.* 5, (2014).



Dot blot analysis of native AAV1-AAV9, AAVrh10 capsids (5E+09-5E+10 capsids) and denatured AAV9 capsids (5E+09-5E+10 capsids, denatured at 95°C for 10 min in sample buffer). The nitrocellulose membrane was blocked with 5% dry milk in PBST (PBS + 0.1% Tween 20) for 1 h at RT. The primary antibody anti-AAV9 (intact particle) mouse monoclonal, ADK9 (Cat. No. 690162) was diluted in blocking buffer (antibody concentration 100 ng/ml) and incubated for 1 h at RT. The secondary antibody goat anti-mouse IgA HRP was also diluted in blocking buffer (antibody concentration 250 ng/ml) and incubated for 1 h at RT. The bands were visualized by chemiluminescent detection using Pierce ECL Plus Western Blotting Substrate.

References

Publication	Species	Application
Hudák, A. et al. Syndecan-4 Mediates the Cellular Entry of Adeno-Associated Virus 9. Int J Mol Sci 24, (2023).	AAV9	FACS
Pan, X., et al., Rational engineering of a functional CpG-free ITR for AAV gene therapy., Gene Ther 29, 333-345, (2022).	AAV9	dot blot
Adachi, K., Enoki, T., Kawano, Y., Veraz, M. & Nakai, H. Drawing a high-resolution functional map of adeno-associated virus capsid by massively parallel sequencing. Nat. Commun. 5, (2014).	AAV9	Neutralization epitope mapping
Varadi, K. et al. Novel random peptide libraries displayed on AAV serotype 9 for selection of endothelial cell-directed gene transfer vectors. Gene Ther. 19, 800-809 (2012).	AAV9	neutralization
Tseng, Y.-S. et al. Generation and characterization of anti-Adeno-associated virus serotype 8 (AAV8) and anti-AAV9 monoclonal antibodies. J. Virol. Methods 236, 105-110 (2016).	AAV9	dot blot, neutralization