

Product datasheet

anti-AAV9 (intact particle) mouse recombinant, ADK9-1R, lyophilized, purified, sample

Short overview

 Cat. No.
 610178S

 Quantity
 10 μg

Concentration 50 μg/ml after reconstitution with 200 μl dist. water

Product description

Host Mouse
Antibody Type Recombinant

IsotypeIgG1CloneADK9-1RImmunogenAAV9 capsids

Formulation Lyophilized; reconstitute in 200 µl dist. water (final solution contains 0.09% sodium azide, 0.5%

BSA in PBS buffer, pH 7.4)

Binding affinity KD value (AAV9) = <1.0E-12 MSynomym Adeno-associated virus 9; AAV-9

Conjugate Unconjugated

Purification Affinity chromatography

Storage before 2-8°C until indicated expiry date

reconstitution

Storage after Up to 3 months at 2-8°C; long term storage in aliquots at -20°C; avoid freeze/thaw cycles

reconstitution

Intended use Research use only

Application Dot blot, ELISA, Neutralization assay

Reactivity AAV9

No reactivity AAV1, AAV2, AAV3, AAV4, AAV5, AAV6, AAV8, AAVDJ, AAVrh10, AAVrh74

Applications

Dot Blot 1:500 (0.1 μg/ml, non-denaturing conditions)

ELISA Assay dependent
Neutralization Assay Assay dependent

Background

For characterization of different stages of infection and very useful for the analysis of the AAV9 assembly process. ADK9-1R specifically reacts with intact AAV9 particles, empty and full capsids. Recognizes a conformational epitope of assembled capsids, not present in denatured capsid proteins and native but unassembled capsid proteins. The antibody cannot be used for immunoblotting. The antibody is also useful for neutralizing experiments. The ADK9-1R antibody recognizes the same epitope as the ADK9 antibody (Cat. No. 690162).

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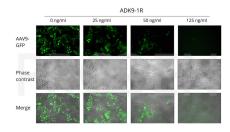
Product images

Antibody clone	Serotype	Binding affinity [KD value]	Neutralization activity [EC50]
ADK9	AAV9	na	~2 ng/ml
ADK9-1R	AAV9	<1.0E-12 M	na
ADK9-h1	AAV9	7.0E-11 M	~8 ng/ml

Comparison of binding affinity and neutralization activity of anti-AAV9 antibodies (mouse monoclonal ADK9, mouse recombinant ADK9-1R and human chimeric ADK9-h1).



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 recombinant, ADK9-1R (Cat. No. 610178) was diluted in blocking buffer (antibody concentration 100 ng/ml) and incubated for 1 h at RT. The secondary antibody goat anti-mouse IgG HRP was also diluted in blocking buffer (antibody concentration 200 ng/ml) and incubated for 1 h at RT. The bands were visualized by chemiluminescent detection using Pierce ECL Plus Western Blotting Substrate.



Neutralization of AAV9-GFP vectors with the ADK9-1R antibody (Cat. No. 610178). AAV infection was shown in HeLa cells and photos (GFP, CPE, merge) were taken ~48 h post infection. Neutralization was enhanced with increasing ADK9-1R concentration.

References

Publication	Species	Application
Haar, J., Blazevic, D., Strobel, B., Kreuz, S. & Michelfelder, S.	AAV9	IA
MSD-based assays facilitate a rapid and quantitative		
serostatus profiling for the presence of anti-AAV antibodies.		
Mol. Ther Methods Clin. Dev. 25, 360–369 (2022).		