

# Datasheet for ABIN7539381

# **AAV8 ELISA Kit**





### Overview

Quantity:	96 tests
Target:	AAV8
Reactivity:	Adeno-Associated Virus 8 (AAV-8)
Method Type:	Sandwich ELISA
Detection Range:	22000000 capsids/mL - 1410000000 capsids/mL
Minimum Detection Limit:	11000000 capsids/mL
Application:	ELISA

#### Product Details

Product Details		
Purpose:	This kit uses sandwich ELISA to determine the titer of AAV capsids in the test sample.	
Sample Type:	Cell Culture Supernatant, Cell Lysate	
Analytical Method:	Quantitative	
Detection Method:	Colorimetric	
Specificity:	Intact AAV wild type virions, AAV recombinant virions, Assembled AAV virions, Intact empty AAV capsids	
Sensitivity:	11000000 capsids/mL	
Components:	<ul> <li>Pre-coated 96-well plate</li> <li>Reference Standard</li> <li>Detection Antibody (20x)</li> <li>HRP Conjugate</li> </ul>	

• Assay Buffer (20x)

- TMB Substrate
- · Stop Solution
- · Instruction manual

# Target Details

Target:	AAV8
Alternative Name:	AAV8 (AAV8 Products)
Application Details	
Sample Volume:	100 μL
Assay Time:	2 h
Plate:	Pre-coated
Protocol:	1. Prepare all reagents, samples and standards,
	2. Wash plate with 1X Assay buffer
	3. Add 100µL standard or sample to each well. Incubate 1 hour at 37 °C,
	4. Aspirate and wash with 300 $\mu$ L 1x Assay buffer 3 times,
	5. Add 100µL Detection Antibody. Incubate 1 hour at 37 °C,
	6. Aspirate and wash with 300 $\mu$ L 1x Assay buffer 3 times,
	7. Add 100µL Streptavidin-HRP conjugate. Incubate 1 hour at 37 °C,
	8. Aspirate and wash with 300 $\mu$ L 1x Assay buffer 4 times,
	9. Add 100 $\mu$ L TMB Substrate. Incubate 10 minutes in the dark at 37 °C,
	10. Add 50μL Stop Solution. Read at 450nm immediately.
Reagent Preparation:	1. Prepare 1X Assay Buffer (Dilute 20X Assay Buffer with distilled H2O)
	2. Prepare 1X Detection Antibody (Dilute the detection antibody (20X) with 1X Assay Buffer)
	3. Preparation of AAV capsid standards: Add 700µL of distilled H2O to one vial of standard.
	Dissolve at room temperature for 10-20 min, mix gently and avoid vortexing. Two-fold serial
	dilution of the AAV8 standard with 1X Assay Buffer for the titration curve in duplicates is recommended.
	4. Sample preparation: Dilute the sample with 1X Assay Buffer to make the capsid titer
	quantification fall within the linear range.
Sample Preparation:	It is recommended to use fresh samples without long storage, otherwise protein degradation
	and denaturation may occur in these samples, leading to false results. Samples should
	therefore be stored for a short period at 2 - 8 °C or aliquoted at -20 °C (≤1 month) or -80 °C (≤
	3 months). Repeated freeze-thaw cycles should be avoided. Prior to assay, the frozen
	samples should be slowly thawed and centrifuged to remove precipitates.
	<ul> <li>If the sample type is not specified in the instructions, a preliminary test is necessary to</li> </ul>

determine compatibility with the kit.

- If a lysis buffer is used to prepare tissue homogenates or cell culture supernatant, there is a
  possibility of causing a deviation due to the introduced chemical substance. The
  recommended dilution factor is for reference only.
- Please estimate the concentration of the samples before performing the test. If the values
  are not in the range of the standard curve, the optimal sample dilution for the particular
  experiment has to be determined.

Assay Precision:

Retest any samples with CV > 10%.

Restrictions:

For Research Use only

#### Handling

Storage:	4 °C/-20 °C

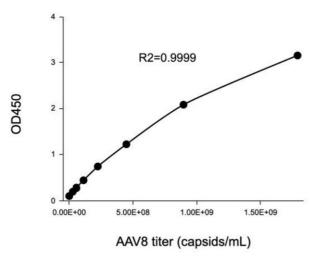
Storage Comment:

Store kit components at 2-8°C. The unopened kit is valid for 12 months from the production date. Reconstituted AAV standard stable at 2-8°C for 2 weeks. -20°C or below for long-term storage and avoid repeated freeze-thaw cycles.

Expiry Date:

6 months

### **Images**



#### **ELISA**

Image 1. Typical standard curve