

## Lassa Virus Pseudovirus

Catalog #: 1007-PP-001

### Description:

The Lassa pseudovirus is a third-generation lentiviral vector pseudotyped with the glycoprotein from the Lassa virus strain Josiah (1976). It contains a firefly luciferase reporter system for sensitive quantification of viral entry.

This pseudovirus is classified under biosafety level 2 (BSL- 2) and is not known or suspected to contain any replicative biological agents due to multiple safety modifications in the viral genome.

**Formulation & Storage:** Supplied in DMEM supplemented with 10% Fetal Bovine Serum. Store at  $-80^{\circ}\text{C}$ . Avoid Thaw/freeze cycle.

### Notes & Usage Guidelines:

Recommended Dilution: 1:50 to 1:100

Luciferase Units for Assays: 300,000 to 600,000

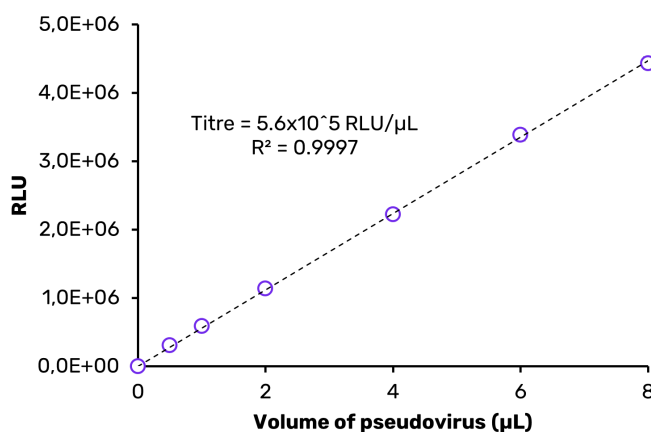
Reporter: Luciferase

~No. of plates that can be tested with a single aliquot: 1 plate

## Applications

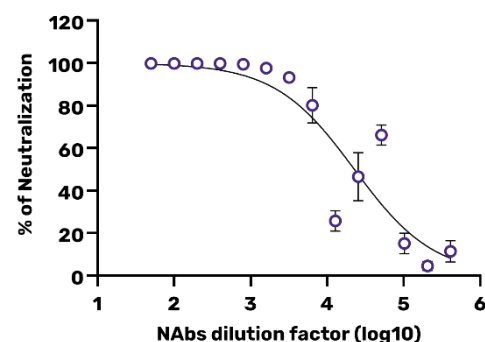
### Viral Titration:

A volume range of pseudovirus (0 – 0.5 – 1 – 2 – 4 – 6 – 8 – 10  $\mu\text{L}$ /well) was mixed in a final volume of 50  $\mu\text{L}$  of medium, in a 96-well plate. Then, 50  $\mu\text{L}$  of medium containing 10,000 cells was seeded in each well. On the day of analysis, an additional 100  $\mu\text{L}$  of Bright-Glo Luciferase reagent was added in each well and incubated for 2 minutes. Data in relative unit luminescence (RLU) were obtained from the analysis of 150  $\mu\text{L}$  of the cell lysate, using a microplate reader. Data are expressed in relative unit luminescence (RLU) as shown below.



### Pseudovirus Neutralization Assay Protocol:

A monoclonal neutralizing antibody (Ab01102-10.0), at a starting dilution of 1/50, was serially diluted in a final volume of 50  $\mu\text{L}$  of complete medium and incubated for 1 hour at  $37^{\circ}\text{C}$ , with 1  $\mu\text{L}$  of pseudovirus, in a 96-well plate. Then, an additional 50  $\mu\text{L}$  containing 10,000 cells was seeded in each well and incubated for 72 hours. Finally, an additional 100  $\mu\text{L}$  of Bright-Glo™ Luciferase buffer was added in each well and incubated for 2 minutes. Data in relative unit luminescence (RLU) were obtained from the analysis of 150  $\mu\text{L}$  of the cell lysate with a microplate reader. Raw data were analyzed using a log(inhibitor) vs normalized-response (variable slope) non-linear regression model in Prism v10 (GraphPad). Percentages of neutralization were normalized considering only cells into wells as 100% neutralization and cells transduced by pseudoviruses without any NABs as 0% neutralization. Data are representative of (duplicates) as shown below.



### Certificate of analysis:

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