SARS-1 Pseudoviral Particles

CATALOG NUMBER: SCV1-PsV-003, 5 mL

Description

Our SARS-1 Pseudoviral Particles are replication-deficient MLV pseudotyped with the SARS-1 spike protein (Genbank Accession # AY278487). They also contain the ORF for firefly luciferase as a reporter. They establish a pseudovirus entry assay for SARS-1 as the spike protein mediated cell entry can be conveniently measured via the luciferase reporter activity. This pseudovirus assay isolates the SARS-1 viral entry from other steps of the viral infection cycle.

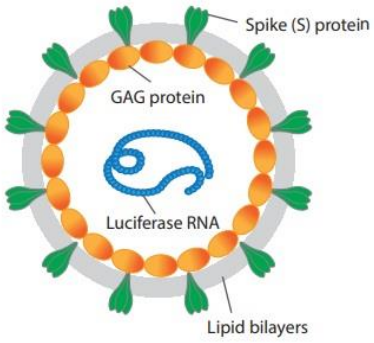

Figure 1. Illustration of the replication-deficient MLV particle pseudotyped with SARS-1 Spike protein

Reference: Identifying SARS-CoV-2 entry inhibitors through drug repurposing screens of SARS-S and MERS-S pseudotyped particles. https://pubs.acs.org/doi/pdf/10.1021/acsptsci.0c00112

Applications

Our Pseudovirus Particles generate robust chemiluminescent signals in cell assays when coupled with our firefly luciferase assay kit (Catalog # CA-L165), useful for 1) screening potential inhibitor to block SARS-1 entry and viral protein translation; 2) measuring the activity of and screening for neutralizing antibody against SARS-1 (refer to the Neutralization Assay Application Note).

Features

- **Robust:** Excellent signal to noise (basal) ratio
- **Easy to use:** Amenable to HTS format (96-well, 384-well and 1536-well format)

Contents

5 mL, for one multi-well plate; PP per mL > 1.0E+07

Storage

Upon receiving this item, store at -70 °C right away. Thaw* before immediate use.
Note: read the instruction for thawing in the following protocol carefully. Do not aliquot and refreeze.

Shelf Life:

Six months from the date of shipping when stored at -70 °C

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ASSAY PROTOCOL

Note: require a luciferase assay reagent (Catalog # CA-L165)

Cell Infection:

1. Count Vero E6 cells or HEK293-ACE2 cells (Catalog # CL-hACE2-002) to be infected and seed ~20K cells per well into appropriate 96-well plates (50 µl per well) DMEM with 10% HyClone™ FetalClone™ II Serum (no antibiotics) or 5K cells per well into appropriate 384-well plates (15 µl per well).
2. Culture cells overnight to make sure the cells stably adhere to the plates.
3. On the 2nd day, remove media, add 50 µl SARS-1 pseudoviral particles* into each well (12.5 µl for 384-well plate). Spin at 700 rpm for 15 min at 4°C.

   Note: thaw the pseudoviral particles quickly in the room temperature water (< 30 minutes, do not shake) and use right away. Discard the unused portion (do not re-freeze or leave it on ice for later use).
4. Incubate for 2 hrs at 37 °C.
5. Add 50 µl DMEM with 10% FC into each well (12.5 µl for 384-well plates).
6. Incubate for 48 hrs at 37 °C.

Measurement of Luciferase Activity in Infected Cells

1. Do not remove medium. Add 100 µl eEnzyme's luciferase assay WORKING SOLUTION (25 µl for 384-well plate) directly into each well. Refer to the protocol of "Firefly Luciferase Assay Kit" (eEnzyme Cat.# CA-L165).
2. Read in a luminescence plate reader and record the data.

Data Analysis

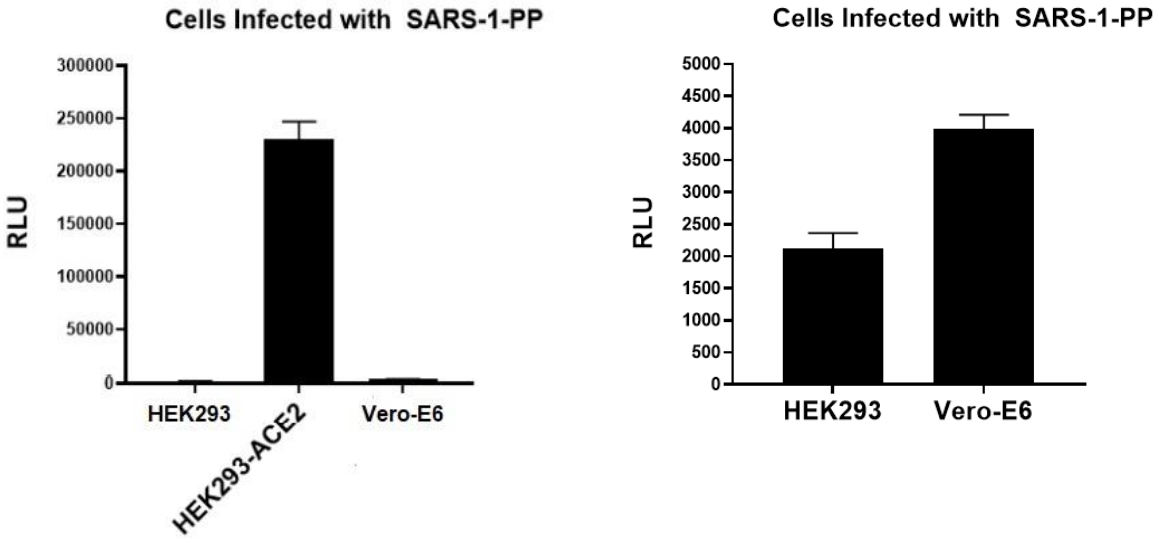

Figure 2. Pseudoviral Particle (PP) Infection Assays
SARS-1 pseudoviral particles with HEK293 cells, or HEK293-ACE2 cells, or Vero-E6 cells in 96-well format.
Legend: SARS-1-PP: MLV w/ SARS-1 spike protein (SCV1-PsV-003)

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