



# Pseudotyped VSV-EBOV Glycoprotein-ΔG-Luciferase (PSVCD101)

This product is for research use only and is not intended for diagnostic use.

## PRODUCT INFORMATION

<b>Product Overview</b>	Recombinant Vesicular Stomatitis Virus pseudotyped Ebola glycoprotein (rVSV pseudotyped EBOV GP) system in which the G protein of VSV has been deleted, replaced with firefly luciferase and used to produce VSV pseudotypes containing the envelope glycoprotein of Ebola virus. Since the infectivity of rVSV pseudotyped EBOV GP is restricted to a single round of replication, analyses of viral entry can be performed using just biosafety level 2 (BSL-2) containment. Infectivity and neutralization of infectivity can be measured by luciferase activity.
<b>Antigen Description</b>	Ebola glycoprotein
<b>Species</b>	Ebola Virus
<b>Concentration</b>	1.97E+08 RLU/ml
<b>Size</b>	20 µl
<b>Buffer</b>	DMEM, 1% FBS, L-glutamine and Penicillin/Streptomycin
<b>Storage</b>	Store at -80°C . Multiple freeze/thaw cycles not recommended. When using the virus, transfer the virus from the -80°C refrigerator and melt it in an ice bath.
<b>Ship</b>	Frozen on dry ice

## BACKGROUND

<b>Keywords</b>	EBOV; Ebola glycoprotein; Ebola virus; Ebola virus glycoprotein; EBOV GP; Ebola virus Pseudovirus; EBOV Pseudovirus; Pseudovirus
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## GENE INFORMATION

## References

1. Whitt, M.A., Generation of VSV pseudotypes using recombinant DeltaGVSV for studies on virus entry, identification of entry inhibitors, and immune responses to vaccines. J. Virol. Methods, 2010. 169(2): p. 365-74.
  2. Howell, K.A., et al., Cooperativity Enables Non-neutralizing Antibodies to Neutralize Ebolavirus. Cell Reports, 2017. 19(2): p. 413-424.
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