

DESCRIPTION

Product	RVP-1426L Bundibugyo ebolavirus (BDBV) Reporter Virus Particles (RVPs)
Lot	FL-1203B
Strain	Democratic_Republic_of_the_Congo/PP_006XHL9.2/2026-05-07
Reporter	<i>Renilla</i> Luciferase
Size	1.0 mL/vial
Packaging	20% FBS/DMEM
Recommended Starting Input	2.5µL per well (96-well plate) for a S:B ≥ 200*
Mycoplasma Test	Negative
Manufacture Date	August 2026
Expiration Date	August 2028

SAFETY & HANDLING

Shipping	Shipped on dry ice
Stability and Storage	Store at ≤ -80°C upon receipt

BDBV RVPs are used to test the ability of serum, antibodies, and drugs to neutralize BDBV infectivity. RVPs display antigenically correct glycoprotein pseudotyped on replication-incompetent virus particles that contain a heterologous VSV core. RVPs are capable of a single round of infection and carry a genome that expresses a *Renilla* luciferase reporter gene upon infection. RVPs are produced in HEK-293T cells using plasmids encoding the glycoproteins.

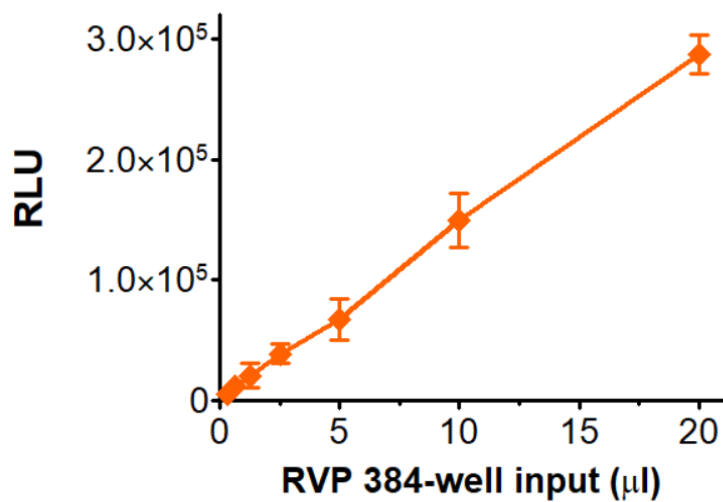
BDBV RVPs are derived from recombinant vesicular stomatitis virus (VSV) from which the VSV glycoprotein gene has been deleted and replaced with a reporter gene (VSVΔG). RVPs are derived from biological materials and should be handled with caution within a BSL2 or enhanced BSL2 laboratory environment. RVPs are not to be used in humans or in animals raised for food.

Thaw tubes in a 37°C water bath for 3 minutes and place on ice until ready to use. RVPs will appear as a translucent, pink solution. Gently mix prior to use and pulse tube for 3 seconds at high speed in a tabletop microfuge to recover all volume from the tube. Vortexing of RVPs should be avoided. Re-freezing of RVPs is not recommended.

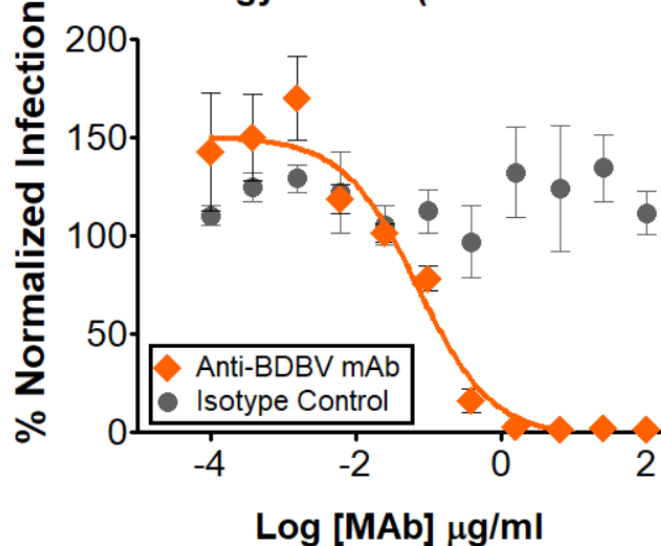
* Determined in the HEK-293T cell line. Input should be optimized based on specific assay conditions and experimental needs. Signal-to-background values may vary depending on instrument make, model, and/or calibration.

INFECTIVITY & NEUTRALIZATION DATA

Bundibugyo RVPs (Lot # FL-1203B)



Bundibugyo RVPs (Lot # FL-1203B)



Infectivity determined in HEK-293T cells (recommended product: ATCC Catalog # CRL-3216). Infectivity values represent the average of three independent vials, with each vial tested in technical quadruplicate.

Neutralization utilized 5 μl of BDBV RVPs in a 384-well plate.

Renilla luciferase activity measured using the Promega *Renilla* luciferase assay system (Promega #E2710). Sample luminescence was read using a PerkinElmer EnVision plate reader.

SIGNAL TO BACKGROUND

RVP 384-well Input (μL)	Signal: Background
20	9,305
10	4,845
5	2,175
2.5	1,260
1.25	677
0.63	398
0.31	186

Signal to background is calculated using mock infected cells as the negative control value.



Christina Go, PhD

Senior Application Scientist, Virology

cgo@integralmolecular.com

Integral Molecular

One uCity Square

25 N. 38th Street, Suite 800

Philadelphia, PA 19104



Jennifer Kilgore

Senior Project Manager/Quality Assurance, Virology

jkilgore@integralmolecular.com

Integral Molecular

One uCity Square

25 N. 38th Street, Suite 800

Philadelphia, PA 19104