



**IBT BIOSERVICES**

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## Rabbit anti-ZEBOV L- Polymerase

**Catalog #:** 0301-045

**Lot #:** 1205002

**Immunogen:** Peptide sequence specific to Zaire Ebola virus (ZEBOV) L-Polymerase protein.

**Description:** Affinity purified rabbit polyclonal antibody reactive to Zaire Ebola virus L-Polymerase. The antibody detects L-Polymerase by Western blot in ZEBOV infected HeLa cells.

**Supplied:** 500 µg of affinity purified antibody is provided in PBS at a concentration of 0.93 mg/mL. 0.01% Sodium azide has been added.

**Raised in:** Rabbits

**Purification:** Antibody is affinity purified using immobilized immunogen.

**Clonality:** Polyclonal

**Relevance:** The antibody can be used for detection of ZEBOV L-Polymerase

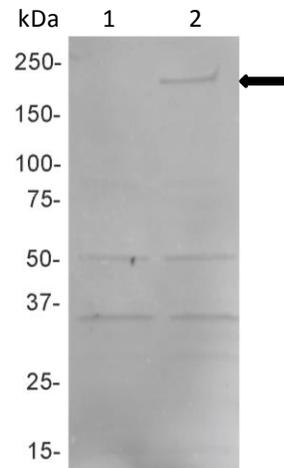
**Storage:** 2-3 weeks +4°C, -20°C long term. Avoid multiple freeze thaws.

**Specificity Testing:**

**ELISA:** Not Tested.

**WB:** Western blot detection was performed to determine the cross-reactivity of the antibody to virus like particles (VLPs) of Zaire and Sudan ebola virus (ZEBOV and SEBOV) and Marburg virus (MARV) expressing glycoprotein, nucleoprotein and VP40. Reactivity to a ZEBOV (Mayinga) ebolavirus infected HeLa cells and uninfected HeLa cells were also evaluated by Western Blot. Cell lysates were prepared in PBS and lysed with RIPA buffer plus protease inhibitors.

### Western Blot Data



Western blot performed with uninfected HeLa cells (Lane 1) and HeLa cells infected with ZEBOV (Mayinga) ebolavirus (Lane 2). Blots were detected using anti-L Polymerase at 1 µg/mL and visualized using an anti-rabbit HRP conjugate and chromogenic substrate (ZEBOV L polymerase is shown by arrow). Data kindly provided by Dr. Olena Shtanko in the lab of Dr. Robert Davey at the Texas Biomedical Research Institute.

**Specificity:** Antibody detects L-polymerase at the expected size of approximately 240 kDa in the HeLa infected lysate (Lane 2), but not the control uninfected lysate (Lanes 1).

**Cross Reactivity:** Western Blot analysis shows that the antibody very weakly detects a band in 2 µg of MARV VLPs between 30-40 kDa but not ZEBOV and SEBOV VLPs. Also, in addition to detection of the expected L-polymerase, the anti-L antibody detects two bands in the uninfected HeLa lysate between 35-52 kDa. The target of this reactivity is unknown. Reactivity of the antibody to L polymerase of other filovirus species is also unknown.

**Intended for research use only, not for human, therapeutic, or diagnostic applications.**

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