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## **Antibody Datasheet**

**Product Name:** Mouse Anti Chikungunya virus VLP Antibody (D7)

**Product:** Monoclonal antibody to Chikungunya virus VLP

Clone Number: D7

**Isotype:** Mouse IgG1, Kappa

Product Code: MAB12385-100

MAB12385-500

**Batch Number:** 

Amount: 0.1 ml

**Concentration:** 1 mg/ml

Physical State: Liquid

**Buffer:** PBS with 0.02% Proclin 300

**Immunogen:** This antibody was raised by immunizing mice.

**Purification:** Protein A affinity chromatography

**Specificity:** This antibody recognises Chikungunya virus-like particles (CHIKV VLPs).

Antibody cross-reacts in ELISA with O'nyong'nyong virus (ONNV) VLPs. It does not cross-react with CHIKV envelope proteins (E1 and E2), Mayaro virus (MAYV) VLPs, MAYV E2 or Ross River virus (RRV) VLPs. See data below.

**Applications:** ELISA, WB

**Secondary Reagents:** Goat anti mouse IgG:HRP (PAB21441HRP)

PanBlock ELISA Blocking Buffer (BUF81201)



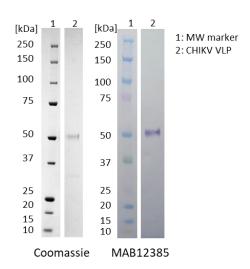


Storage:

Store at 4°C for up to 3 months. For longer storage, aliquot and store at -20°C. Avoid cycles of freezing and thawing. The antibody is shipped at ambient temperature.

## **Western Blot**

100ng of antigen was used for SDS-PAGE, in reduced form. Proteins were transferred using Transblot for 7 min. at 25V. 5% dry milk in PBS-T was used as blocking buffer and dilution buffer for antibodies. Primary antibodies and goat anti-mouse-IgG-HRP secondary antibody (Biorad 103005) were used at 1:1000. All steps were carried out for 1h at room temperature with gentle rocking. KPL Membrane TMB was used for detection. Development time 30 sec.



## **ELISA**

Coat plate with  $0.5\mu g/ml$  of antigens in 1X DPBS overnight at 2-8 degrees. Block for 1.5 h using 1%BSA/DPBS.300 $\mu$ l/well. Wash plate 3X using Tris wash buffer. Add 100 $\mu$ l of prepared antibody dilutions. Incubate for 2 h at RT at 800rpm. Wash plate 3X using Tris wash buffer. Add 100 $\mu$ l of Goat anti mouse IgG HRP antibody (1:5000 dilution) to all the wells and incubate for 1 h at RT,800rpm. Wash plate 4X using Tris wash buffer. Add 100 $\mu$ l HKTMB and incubate for ~2 min. at RT. Add 100 $\mu$ l 1M HCl to stop the reaction and read the plate at 450nm.

