

ECHOVIRUS TYPE 6 Purified Viral Lysate

PRODUCT DESCRIPTION:

Echovirus is a non-enveloped enterovirus that contains a linear, positive-sense single-stranded RNA.

Echovirus Type 6 is propagated in the Vero cell line.

The virus is purified using sucrose density gradient ultracentrifugation, disrupted in the presence of 0.5% Triton X-100 non-ionic detergent/0.6 M KCl, and heat inactivated.

Echovirus lysate is sold in vials containing 1.0 mg of protein, and is shipped on dry ice. Protein concentrations generally range from 0.5 to 3.0 mg/ml.

Custom orders are available, including specific buffer formulations and package sizes.

INTENDED USE:

This product is intended for research, product development, quality assurance testing, or further manufacturing use.

Viral lysates can be utilized as an antigen, as a source for the purification of viral proteins, or for the detection of viral antibodies. Applications include:

- Immunodetection of antibodies to Echovirus using solid-phase enzyme immunoassays (EIA)
- Western blot
- Dot blot
- Other protein-based assays

This product was manufactured in a facility which has a Quality Management System that is certified as being in compliance with ISO 13485.

ETIOLOGIC STATUS/BIOHAZARD TESTING:

Echovirus is a Biosafety Level 2 organism.

Viral inactivation is verified for every lot of lysate by the absence of viral growth in validated tissue culture based infectivity assays.

PRECAUTIONS:

USE UNIVERSAL PRECAUTIONS when handling this product!

Lysate has been treated by a method validated to be effective for virus inactivation. However, no method can be guaranteed 100% effective.

This material should be handled as if capable of transmitting infectious agents.

RECOMMENDED STORAGE:

The viral lysate is stable for at least one year when stored at -65°C or below.

To avoid repeat freeze-thaws, which could negatively impact product performance, viral lysate should be stored in aliquots upon receipt.

DO NOT USE IN HUMANS!

These products are NOT intended for use in the manufacture or processing of injectable products subject to licensure under section 351 of the Public Health Service Act, or for any other product intended for administration to humans.

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