ZeptoMetrix[®]

Catalog Number: 0810229CFHI

Coronavirus (CoV) Strain: 229E Culture Fluid (Heat Inactivated) (1 mL)

PRODUCT DESCRIPTION:

Coronavirus (CoV) (Strain: 229E), a Group 1 CoV, is an enveloped virus with spiked projections, and contains a nucleocapsid with positive-sense RNA.

Each frozen aliquot contains 1 mL of heat inactivated viral culture fluid. The pre-inactivation titer was determined from an infectious aliquot.

Viral inactivation is verified after heat inactivation by the absence of viral growth in tissue culture-based infectivity assays.

INTENDED USE:

Heat inactivated viral culture fluids are sold as consumable testing materials. The suitability and performance characteristics should be determined by your laboratory for each intended usage.

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.

FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES.

BIOSAFETY:

Please consult your institution's regulations regarding the use of this product. For a detailed discussion on biological safety, see the 5th edition of Biosafety in Microbiological and Biomedical Laboratories (BMBL), published by the CDC at

http://www.cdc.gov/biosafety/publications/bmbl5/inde x.htm

PRECAUTIONS:

- Use Universal Precautions, this product is potentially biohazardous.
- Repetitive freezing and thawing is not recommended (aliquot material if necessary).

To avoid cross-contamination, use separate pipette tips for all reagents.

RECOMMENDED STORAGE:

Heat inactivated viral culture fluids should be stored at -65°C or below.

PI0810229CFHI Revision: 03 Effective Date: 10/26/2021

REF	Catalog Number	X	Temperature Limitation
LOT	Batch Code	M	Expiration Date
RUO	For Research Use Only	8	Biological Risk
	Manufacturer		

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www.ZeptoMetrix.com ZeptoMetrix LLC • 878 Main Street, Buffalo, NY 14202 USA • Tel (800) 274-5487 This product was manufactured in a facility which has a Quality Management System that is ISO 13485 certified.