Catalog Number: **NATTFP-BIO**

NATtrol™ Tropical Fever Verification Panel

PRODUCT DESCRIPTION:

NATtrol™ Tropical Fever Verification Panel (NATTFP-BIO) contains purified, intact bacteria and viruses that have been chemically modified to render them non-infectious and refrigerator stable. The panel members are supplied in a proprietary matrix.

INTENDED USE:

NATtrol™ Tropical Fever Verification Panel can be used to develop procedures for verification of performance of molecular assays that detect the presence of bacterial, protozoan and viral nucleic acids (from organisms listed in Table 1). The panel can also be used for training and evaluation of operator proficiency.

MATERIALS SUPPLIED:

9 x 0.3 mL vials (positive panel members) and 3 x 1.0 mL vials (negative control) as listed in Table 1.

WARNINGS AND PRECAUTIONS:

- NATtrol™ inactivation was carried out on virus and microorganism stocks used to formulate the panel members. The inactivation was verified in a standard microbiological growth protocol.
- This product contains inactivated viruses and microorganisms and materials of human and animal origin. Safe practices suggest that the product be considered potentially infectious and to use Universal Precautions when handling.
- Refer to CDC guidelines and local regulations for handling and disposal.
- The matrix used in the manufacture of this product is treated with 0.09% sodium azide. It was manufactured from Human Serum Albumin that has been tested and found to be non-reactive at the donor level for HIV-1/HIV-2 Antibody, HBsAg and HCV Antibody by FDA licensed donor screening test methods. All materials are also tested for HIV-1 and HCV by FDA approved Nucleic Acid Test (NAT) methods.
- Heat inactivated Fetal Bovine Serum used in the manufacture of this product meets applicable USDA requirements for abattoir sourced animals, traceability, and country of origin. The materials were collected at USDA licensed establishments or legally imported from countries recognized by the USDA as negligible or controlled for risk for Bovine Spongiform Encephalopathy (BSE) and other exotic disease agents. Donor animals were inspected ante and postmortem at the abattoir as required by the USDA.
- Do not use past the expiration date on the label.
- To avoid cross-contamination, use separate pipette tips for all materials.

RECOMMENDED STORAGE:

Store at 2-8°C.

INSTRUCTIONS FOR USE:

- Mix tube vigorously for at least 5 secs.
- Process according to manufacturer's instructions for sample to result assays.
- Extract nucleic acid prior to use in downstream assays that are not sample to result.
- Each vial is intended for single use.

LIMITATIONS:

- FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES.
- Quality control materials should be used in accordance with local, state, federal, and accreditation requirements.
- This product is not intended to replace the manufacturer's controls provided with the assay.

PINATTFP-BIO Revision: 00

Effective Date: 02/18/2025



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EXPECTED RESULTS:

Each laboratory must evaluate the panel and establish their own performance criteria using their specific assay. NATtrol™ Tropical Fever Verification Panel does not have assigned analyte values or ranges.

TABLE 1: PANEL MEMBERS

Panel Member (Strain)				
Chikungunya Virus (R80422)				
Dengue Virus Type 1 (Hawaii)				
Dengue Virus Type 2 (New Guinea C)				
Dengue Virus Type 3 (H87)				
Dengue Virus Type 4 (H241)				
E. coli with Dengue Virus Type 2 (Dak Ar A1247) plasmid - recombinant ¹				
E. coli with Leptospira plasmid - recombinant1				
E. coli with P. falciparum/vivax plasmid - recombinant1				
E. coli with P. spp/ovale plasmid - recombinant1				
Negative Control				

¹This analyte only contains a short sequence of the organism's genome therefore each laboratory must evaluate performance in their assay.

Symbols used in the labeling of this product:

	Manufacturer	1	Temperature Limitation
RUO	For Research Use Only	Ω	Expiration Date
REF	Catalog Number	₩	Biological Risk
LOT	Batch Code		

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