

Malaria P. falciparum Culture Panel **Product Code: KZMC043** Strain: W2 (Type B)

Lot Number: 1611-272-00023

For Research Use Only. Not for use in diagnostic procedures.

NAME AND INTENDED USE:

The Malaria P. falciparum Culture Panel is intended for use with in vitro tests for the determination of the presence or absence of P. falciparum based on the detection of the histidine rich protein 2 (HRP2) or assays for the quantification of HRP2. This panel is for Research Use Only and should not be used in diagnostic procedures.

SUMMARY:

The Malaria P. falciparum Culture Panel can be used to evaluate the analytical performance of HRP2 detecting assays. Each panel contains aliquots of a P. falciparum laboratory strain at seven predetermined HRP2 concentrations as well as negative aliquots. This panel can be used to evaluate the performance of assay or diagnostic test based on the detection or quantification of HRP2.

PRINCIPLES OF THE PROCEDURE:

The Malaria P. falciparum Culture Panel has been designed for use with in vitro test for detecting the presence or absence of P. falciparum through the detection of HRP2 and with laboratory assay for the quantification of HRP2. The Malaria P. falciparum Culture Panel is prepared from human and nonhuman components. The Malaria P. falciparum Culture Panel reagents should be evaluated as unknown specimens per the instructions supplied by the manufacturer of the in vitro assay being used.

REAGENTS:

The Malaria P. falciparum Culture Panel reagents contain culture-derived P. falciparum parasites diluted in human whole blood at decreasing HRP2 concentrations as follows:

- 1. Two vials 8,000 pg/mL (200 µL per vial).
- 2. Two vials 800 pg/mL (200 µL per vial).
- 3. Two vials 400 pg/mL (200 µL per vial).
- 4. Two vials 120 pg/mL (200 µL per vial).
- 5. Two vials 40 pg/mL (200 μ L per vial). 6. Two vials 12 pg/mL (200 μ L per vial).
- Two vials 4 pg/mL (200 µL per vial).
- 8. Two vials Negative (200 µL per vial).

WARNINGS AND PRECAUTIONS:

- FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES.
- USE UNIVERSAL PRECAUTIONS: HANDLE AS IF CAPABLE OF TRANSMITTING INFECTIOUS AGENTS.

The Malaria P. falciparum Culture Panel reagents are prepared from P. falciparum laboratory culture samples and human whole blood. All materials used in the preparation of The Malaria P. falciparum Culture Panel reagents are known to be non-reactive for HIV1/2 Ab, HBsAg and HCV Ab. However, no known test method can assure that products derived from human sources will not transmit infection. It is recommended that these reagents and all human specimens be handled in accordance with Biosafety Level 2 practices as described in the CDC NIH publication, Biosafety in Microbiological and Biomedical Laboratories (1), or other equivalent guidelines (2,3).

SAFETY PRECAUTIONS:

- 1. Clean any spillage immediately and thoroughly using a suitable disinfectant such as 1% bleach solution.
- 2. Handle and dispose of all specimens, controls and materials used in testing as though they contain infectious agents (1-3).

HANDLING PRECAUTIONS:

- 1. Do not use Malaria P. falciparum Culture Panel reagents beyond the expiration date.
- 2. Avoid contamination of reagents when opening and dispensing.

STORAGE INSTRUCTIONS:

- 1. Store Malaria P. falciparum Culture Panel reagents long term (≥30 days) at ≤ -70°C or Short Term (≤30 days) at ≤ -10°C when not in use.
- 2. Vials should be stored upright to prevent leakage.
- 3. For single use only.

INDICATIONS OF REAGENT INSTABILITY OR DETERIORATION:

1. Alterations in physical appearance may indicate instability or deterioration of Malaria P. falciparum Culture Panel.

PROCEDURE:

- 1. The Malaria P. falciparum Culture Panel reagents may be included in a test run following the procedure provided by the in vitro assay manufacturer for unknown specimens.
- 2. Allow The Malaria P. falciparum Culture Panel reagents to thaw on ice prior
- 3. Mix contents by gentle swirling prior to use. Do not mix by vigorous shaking.
- 4. Vial should be considered for single use only. The HRP2 concentration of vials undergoing several freeze-thaw cycles cannot be ensured.

INTERPRETATION OF RESULTS:

The Malaria P. falciparum Culture Panel test results should be determined as recommended for unknown specimens in the package insert for each commercially available test kit and compared to the reagent predetermined HRP2 concentration.

LIMITATION OF THE PROCEDURE:

- 1. Malaria P. falciparum Culture Panel reagents must not be substituted for the positive and negative control reagents provided with some commercially available test kits.
- 2. Malaria P. falciparum Culture Panel reagents are provided for Research Use Only and should not be used in diagnostic procedures, for calibration or as primary reference preparations for any test kit.
- 3. PROCEDURE and INTERPRETATION OF RESULTS provided in package insert of each commercially available test kit must be followed closely when testing the Malaria P. falciparum Culture Panel reagents. Deviations from the recommended procedures may produce unreliable results.
- 4. It is the responsibility of each laboratory to determine the suitability of Malaria P. falciparum Culture Panel reagents for its particular use. They also must establish guidelines for the interpretation of results.

SPECIFIC PERFORMANCE CHARACTERISTICS:

The Malaria P. falciparum Culture Panel reagents were tested using commercially available test systems following the procedures provided by the manufacturer for the testing of unknown specimens. The data contained in the Panel Box Key Code is intended to be for informational use only. Each laboratory should establish its own performance characteristic.

This Product was manufactured in a facility which has a Quality Management System that is ISO 13485 certified. ZeptoMetrix Corporation • 25 Kenwood Circle, Franklin, MA 02038 • Tel (508) 553-5800 • Fax (508) 520-1525

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Product Key Code Expiration Date: 01/06/2021

Variable	Values	Description	Comment
STUDY ID	01	Malaria specimen repository	Constant
CULUTRE PRODUCTION SITE	06	Supplier	N/A
STRAIN AND [HRP2]	0032	W2, 8000 pg/mL	Type A
	0038	W2, 800 pg/mL	Type A
	0001	W2, 400 pg/mL	Type A
	0002	W2, 120 pg/mL	Type A
	0003	W2, 40 pg/mL	Type A
	0004	W2, 12 pg/mL	Type A
	0005	W2, 4 pg/mL	Type A
	0031	Negative, n/a pg/mL	N/A
TYPE OF SPECIMEN	2	Culture specimen	Constant
ALIQUOT ID	001-999	Unique identification for each aliquot	

This data is intended to be for informational purposes only. They are not intended to represent performance specifications

REFERENCES:

- 1. U.S. Department of Health and Human Services. Biosafety in Microbiological and Biomedical Laboratories. HHS Publication (NIH) 93-8395. Washington: U.S. Government Printing Office, May, 1993.
- 2. National Committee for Clinical Laboratory Standards. Protection of Laboratory Workers from Infectious Disease Transmitted by Blood, Body Fluids and Tissue Second Edition, Tentative Guideline. NCCLS Document M29-T2. Villanova, PA: NCCLS, 1991.
- 3. National Committee for Clinical Laboratory Standards. Clinical Laboratory Waste management; Approved Guideline> NCCLS Document GP 5-A. Villanova, PA: NCCLS, 1993.

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