Instructions for Use

Version: 2.1.3

Revision date: 25-Nov-24



Human Hepatitis E Virus IgM (HEV IgM) Rapid Test Kit

Catalog No.: abx092089

Size: 20 tests / 40 tests

Storage: Store all reagents at 4-30°C. Keep dry.

Application: For qualitative detection of Human Hepatitis E Virus IgM in serum and whole blood.

Introduction and assay principle

Abbexa's Human Hepatitis E Virus IgM Rapid Test Kit is based on the gold immuno-chromatography assay (GICA) principle. Any Human Hepatitis E Virus IgM present in the samples combines with the colloidal gold particle-labelled Human Hepatitis E Virus antigen. When the concentration of Human Hepatitis E Virus IgM in the sample is more than the detection limit, there is a color change in the detection line and the result is positive. When the concentration of Human Hepatitis E Virus IgM in the sample solution is less than the detection limit, there is no color change in the detection line and the result is negative.

Kit Components

Test cassettes: 20 / 40

Sample diluent: 1 vial / 2 vials

Material Required But Not Provided

- High-precision pipette and sterile pipette tips
- Timer
- Lancets

Sample preparation

Whole blood samples should be tested immediately after collection or stored at 2-8 °C for up to 24 hours. Serum samples should be collected using conventional methods and stored between 2-8 °C for short-term storage (up to 5 days) or -20 °C for long-term storage. Fresh samples are recommended. Avoid repeated freeze/thaw cycles, bacterial pollution, visible particles; and avoid cloudy, hemolytic, or viscous samples.

Assay procedure

- 1. Take a test cassette and lay it flat on a clean table. Slowly and vertically add 10 µl of sample to the sample well, then add 2 drops (approximately 70-90 µl) of Sample diluent to the sample well on the test cassette. Avoid foaming. Start the timer.
- 2. Results may be read within 15 20 minutes. Results after 20 minutes are not valid, and we recommend discarding test cassettes after 20 minutes to minimize confusion.

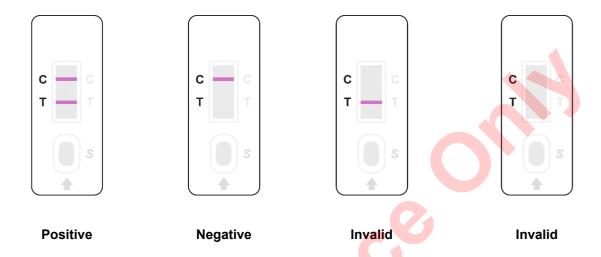
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Results analysis

- Positive result: A colored line is observed in both the control (C) section and the test (T) section.
- Negative result: A colored line is observed in the control (C) section but not the test (T) section.
- Invalid result: No colored line is observed in the control (C) section.



Notes

- 1. The test cassettes should be brought to room temperature before use.
- 2. After opening the aluminum foil, use the test cassette as soon as possible.
- 3. Samples should be clear with no visible particles, turbidity or bacterial pollution.
- 4. Do not mix or re-use the disposable pipettes to avoid cross-contamination.
- 5. Do not use water, PBS, or similar solutions as the negative control.
- Avoid touching the cassette membrane through the sample well or test result window.
- 7. This kit is for qualitative detection of Human Hepatitis E Virus IgM in human serum, plasma and whole blood. For other sample types, a preliminary experiment is recommended to determine compatibility with this kit.
- 8. This kit is for research use only and the results are for reference only. It is recommended to use this kit in conjunction with another detection method.
- 9. All waste should be disposed of appropriately. Please note that you may need to follow special waste disposal procedures for infectious samples. Please check local disposal regulations.