DiAGSure Zika virus Detection Kit

20 Tests

For research use only

DiAGSure DIAGNOSTIC
For the people, by the people

GCC Biotech (India) Pvt. Ltd.
tech.support@gccbiotech.co.in | 91-33-24951044/24950004

Description:

Zika is a viral disease spread by *Aedes* mosquito. It was first identified in Uganda to occur in monkeys wherefrom it spread to humans crossing the species barrier. The disease is now reported to occur globally. Although most cases of zika infection are asymptomatic, the common symptoms generally include mild and include fever, rash, conjunctivitis, muscle and joint pain, malaise or headache. The symptoms typically last for 2-7 days. Zika virus can be transmitted to the fetus from an infected mother that lead to microencephaly and other congenital disorders in the new-born, often referred to as congenital Zika syndrome. The icosahedral virus belongs to Flaviviridae (Baltimore Group IV) and its genome is a ss-(+)-RNA of about 10kb in length encoding structural and non-structural (NS) proteins. Zika virus infection can be diagnosed by laboratory tests of blood and other body fluids like urine or serum. RT-PCR based detection of the virus efficiently detects the presence of viral RNA in infected human blood and other body fluids.

DISCLAIMER: The DiAGSure Zika virus Detection Kit has been designed for *in-vitro* use only.

Intended Use:

This kit amplifies a unique 350-bp sequence specific for Zika virus and is absent in other closely-related Flaviviridae members. This kit also contains a standard marker for size comparison of the amplicon.

Principle:

The DiAGSure Zika virus Detection Kit involves semi-quantitative RT-PCR based detection of a conserved specific 350-bp sequence in the Zika viral genome using gene-specific primers. PCR-based detection is

emerging as a highly sensitive diagnostic tool for the detection of pathogen in a wide array of clinical samples. Reverse transcriptase converts the viral RNA to cDNA which serves as a template for PCR. A basic PCR reaction involves three basic steps:

i. Denaturation, where separation of the two DNA strands occur
ii. Annealing, where the primers are allowed to anneal to their cognate templates
iii. Extension, where the actual amplification occurs that is repeated between 25 and 40 cycles in each assay. The PCR primers have been designed to ensure high specificity and sensitivity.

Features:

- ✓ Fast and simple
- ✓ Rapid detection of the virus in clinical samples
- ✓ Highly sensitive
- ✓ Specific detection of the virus
- ✓ Reproducibility of results

Storage and Shelf life:

The provided kit has a shelf-life of 6 months when stored at -20°C. Repeated thawing and freezing of RT enzyme and PCR reagents may reduce the sensitivity and therefore should be avoided. If reagents are to be used multiple times, we recommend storing reagents as aliquots to avoid repeated freeze and thaw. The degradation of sample RNA specimens may also compromise with the sensitivity of the assay. Usage of the kit after the expiry date stated on pack is not recommended.

Kit contents:

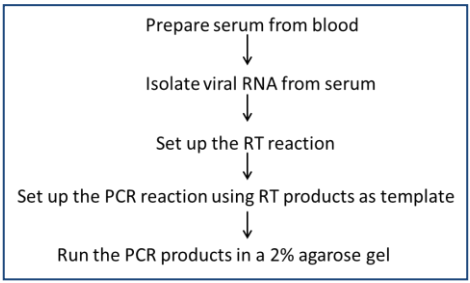
(Storage: -20°C, in a Frost-free freezer):

Kit Contents	Volume for 20 tests
10X RT buffer	50 µL
GRTScript Reverse Transcriptase	25 µL
dNTP-Primer Mix	50 µL
ZiV Primer mix	45 µL
DiAGPol PCR Master Mix	500 µL
DiAGSure DNA ladder	100 µL
Gel loading dye	100 µL
Nuclease free water	500 µL

Sample Material Preparation:

The DiAGSure Zika Detection Kit detects the presence of the Zika virus in human blood samples. Isolate total RNA from blood. Use a specified amount (see below) of this RNA and prepare cDNA which has to be used as a template for amplification of the 350-bp region.

Basic workflow:



Starting volume of serum: 200µL
Elution volume: 30µL

RT reaction set up and conditions:

1. For setting up the RT reaction (10µL total volume), add the following reagents in a 0.2mL PCR tube and mix by pipetting.

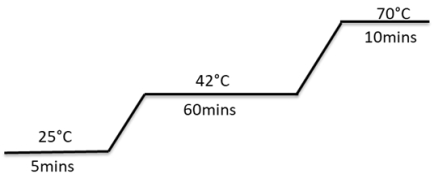
Isolated RNA	6 µL
dNTP-Primer mix	2 µL

2. Incubate the mix at 65°C for 5mins followed by quick-chill on ice for 2-3 mins.
3. Add the following reagents to the tube:

10X RT buffer	1 µL
GRTScript Reverse Transcriptase	1 µL

4. Mix vigorously and pulse spin to bring the contents to the bottom of the tube. Place the tube in a thermal cycler and run the RT reaction for a single cycle under the following cycling conditions:

Stage	Temperature (°C)	Time
Annealing	25	5mins
Extension	42	60mins
Inactivation	70	10mins
Final hold	4	∞



Diagrammatic view of RT reaction conditions

PCR Protocol:

To the RT Reaction tube, add the following constituents:

Template cDNA	10 µL
DiAGPol PCR Master Mix	9 µL
ZiV primer mix	1 µL

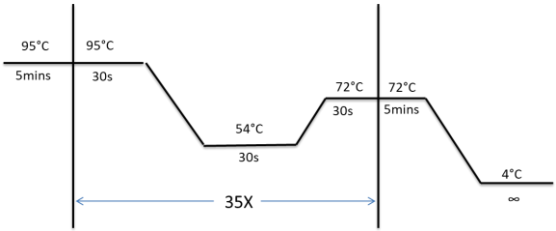
Set up a No Template Control (NTC) reaction, replace cDNA with 10 µL of Nuclease free water and add ZiV primer mix and DiAGPol PCR Master Mix accordingly.

N.B.: If required, the volume of template may be increased accordingly.

Mix vigorously by pipetting up and down and pulse-spin to bring the contents to the bottom of the tube and place the tube in following thermal cycling program.

PCR conditions:

Stage	Temperature (°C)	Time	No. of cycles
Initial denaturation	95	5 mins	1
Denaturation	95	30 secs	35
Annealing	54	30 secs	
Extension	72	30 secs	
Final extension	72	5 mins	1
Final hold	4	∞	1



Diagrammatic view of PCR cycling conditions

Add 1µL of the supplied gel-loading dye to the PCR products, mix well and run the PCR products along with 5µL of the supplied DiAGSure DNA ladder in a 2% agarose-TAE gel.

Results Interpretation:

The presence of a 350-bp band lying between 400bp and 500bp with respect to the standard marker indicates the presence of the Zika virus in the clinical sample. The absence of the 350-bp band in the test sample indicates the absence of Zika virus infection (See Fig 1).

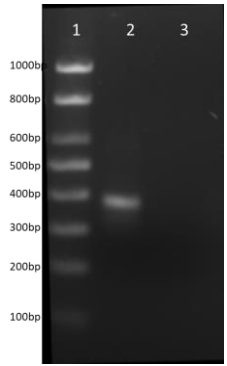


Fig 1. Representative gel image showing amplification of the Zika sequence. Lane 1: DiAGSure DNA ladder; Lane 2: Positive amplification of the Zika amplicon at 142-bp; Lane 3: Negative control.

Sensitivity:

The DiAGSure Zika virus Detection Kit is highly sensitive and can detect up to 3 attomoles of the virus under *in vitro* conditions.

Quality Control:

All reagents in the DiAGSure Zika virus Detection Kit are free from endonuclease and exonuclease activities and the kit has been functionally tested for amplification.

Precautions:

- Ensure that RNA has been properly isolated.
- Use freshly isolated RNA for amplification.
- The working desk for RNA isolation should be clean and properly wiped with 70% ethanol.
- Clean the working area and the nozzles of the pipette with RNaseZIP (Cat. No. G7111; Not provided).
- All microcentrifuge tubes and Pipetman tips should be double-autoclaved.
- The RT reaction should be set up meticulously on ice and carried out under conditions as indicated.
- Verify that all reagents are added to the PCR reaction in indicated amounts and proper PCR conditions have been maintained.
- It is advisable to store the reagents in aliquots for multiple uses.

Safety information:

The DiAGSure Zika virus Detection Kit is for laboratory use only. Use proper safety measures while handling clinical samples, like wearing mask, gloves, lab-coat, etc.

Technical assistance:

Satisfaction of the customers is our utmost priority. For any kind of technical assistance, always feel free to reach out to us at tech.support@gccbiotech.co.in.