

[Product Name] West Nile Virus (WNV) N ic Acid Test Kit (Lyophilize s]

[Package Specificati 16 T/box

[Intended Use] The kit uses fluorescent PCR to detect West Nile Virus (WNV). This product requires the operation of a fluorescent quantitative PCR instrument to achieve rapid detection of POCT.

Institutent to achieve rapid detection OF POCT. **[Testing Principlo]** The test kit uses nucleic acid extraction reagents to extract the nucleic acid (DNA/RNA) from the sample. Under the acid on 6 a high-efficiency reverse transcriptase, cDNA complementary to the RNA template is synthesized in a one-step reaction using RNA as the template. Under the acidon of Tag enzyme, the copy number of the specific target fragment is amplified through cycles of high-temperature denaturation, annealing at a moderate temperature, and extension using DNA as the template. The fluorescence-labeled specific probe hybridizes with the amplified target fragment, and the 55-x3' exonuclease activity of Tag polymerase separates the reporting group and quencher group of the fluorescence probe, emitting a specific fluorescence signal. The specific fluorescence is gial is detected using a fluorescence sample and the formation of the amplification curve.

[Contents]

Item	Quantity	Storage	
PCR master mix	16 pcs	-20°C (Away from light)	
Instructions for use	1 pcs	- Room Temperature	
Sample buffer	16 pcs		
Swab	16 pcs		
Biohazard bag	16 pcs		

[Storage conditions and shelf life] 1. Shelf life: 24 months. 2. Production date and expiration date are on th e pa ckade.

[Compatible Instruments] This test kit is compatible with FLASHTEST r fluorescence PCR instrument.

[Sample] EDTA anticoagulate

d blood

[Sample Handling] EDTA anticoagulated blood: Collect blood using a blood collection tul containing EDTA anticoagulant. 1. Add 100 µL of blood to the sample buffer with a disposable dropper 2. Thoroughly mix the sample buffer with a repetitive pipetting action, using the disposable dropper. 3. Add 200 µL of mixed buffer to the nucleic acid extraction cartridge extraction.

[Specimen storage] Samples used for nucleic acid extraction and detection should be tested as soon as possible. Samples to be tested within 24 hours can be stored at 4°C. Samples that can not be tested within 24 hours should be stored at -20°C for up to 10 days. Avoid repeated freezing and thawing of samples.

Constructions for Use)
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 Instructions for Use)
 Inded 20µL of elution from magnetic bead extraction, to each PCR tube
 Close the lid tightly.
 Is Shake all the liquid to the bottom of the PCR tube. Use the vortex
 mixer to mix the PCR tube PCR tube beschart mixing make
 sure all liquid is at the bottom of the PCR tube, by shaking the tube again
 (optional: use a small centrifuge for 3 seconds to shift all liquids to the
 bottom.)

2. PCR Amplification 2.1 Set the parameters as fol

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Temperature 55°C 94°C Step Time Cycle 3min 30s 1 1 2 1 55

94°C 58°C is 20

Channel	FAM	VIC	ROX	Cy5
Target		Internal reference		WNV

20 s ×40

3. Result Interpretation 3.1 Reference Range:

Parameter	Reference Range	Result Interpretation
Internal Control	Ct ≤ 37 and there is a clear exponential amplification curve	Valid
	Ct > 37 or No Ct	Invalid
Pathogen	Ct ≤ 37 and there is a clear exponential amplification curve	Positive
	Ct > 37 or No Ct	Negative

3.2 Test Result Interpretatio

Pathogen Result	Internal Control Result	Test Result Interpretation	
Positive	Valid	Pathogen Positive	
Negative	Valid	Pathogen Negative	
Any Result	Invalid	Test invalid, please retest	

[Test Limitations] 1. The test results of this kit should be comprehensively analyzed in conjunction with other relevant physical examination results and should not be used as the sole basis for diagnosis. 2. Improper sample collection, transportation, storage, handling, and inadequate laboratory conditions may lead to inaccurate results. 3. Other unconfirmed interferences or PCR inhibitors may lead to false negative results. 4. Sequence variations caused by mutations or other factors in the targe gene of the virus being tested may lead to false negative results.

(Product Performance) I. Positive and negative control consistency: The positive and negative controls included in this test kit have been tested with the company's working reference materials, and the positive and negative compliance rates are both 100%.
3. Sensitivity: Imil of detection is 500 copies/mL.
3. Specificity: This assay does not cross-react with non-target pathoge samples.
4. Precision: The coefficient of variation (CV, %) of the Ct values for 10 consecutive tests of one strong positive sample and one weak positive sample is ≤5%.

Sample 19 source.
[Notes]
1. Before using a PCR kit, check the lyophilized PCR mix at the bottom
of the tube is in good condition (white and clumped). Liquified lyophilized
PCR mix can not be used. After opening, it should be used as soon as
possible or stored away from light.
2. This product is only for in vitro testing (for animals). All operations must
strictly follow the instructions.
3. Overloading samples may result in false negatives. Retest is
recommended.
4. Avoid bubbles in PCR tubes. Keep the tube cap firmly closed.
5. Use disposable lips, gloves, and laboratory coats.
6. After tests, ideinfect the workbench with 10% hypochlorous acid, 75%
ethanol, or UV light.
7. All items in the kit should be treated as biowaste and handled in
accordance with local laboratory regulations.