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## Hantavirus Pulmonary Syndrome Real Time RT-PCR Kit

### User Manual

#### For In Vitro Diagnostic Use Only

**REF** QR-0090-01

For use with LightCycler1.0/2.0 Instrument

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#### 1. Intended Use

Hantavirus pulmonary syndrome real time RT-PCR kit is used for the detection of Hantavirus pulmonary syndrome (HPS) in serum by using real time PCR systems.

#### 2. Principle of Real-Time PCR

The principle of the real-time detection is based on the fluorogenic 5' nuclease assay. During the PCR reaction, the DNA polymerase cleaves the probe at the 5' end and separates the reporter dye from the quencher dye only when the probe hybridizes to the target DNA. This cleavage results in the fluorescent signal generated by the cleaved reporter dye, which is monitored real-time by the PCR detection system. The PCR cycle at which an increase in the fluorescence signal is detected initially is proportional to the amount of the specific PCR product. Monitoring the fluorescence intensities in real-time allows the detection of the accumulating product without having to re-open the reaction tube after the amplification.

#### 3. Product Description

Hantaviral diseases in humans are caused by a group of closely related, trisegmented, negative-sense RNA viruses of the genus Hantaanvirus, of the family Bunyaviridae. The type and severity of the disease depends largely on the serotype of the virus involved. Two classes of hantavirus-associated illnesses have been described: HFRS for the disease in which the kidneys are primarily involved, and HPS for the disease in which the lungs are primarily affected. The recent data concerning the pathogenesis of Hantavirus infection (e.g. Sherif R. Zaki) confirm that the basic necroinflammatory changes in the infection develop in the blood vessels. The endothelium is the internal cover of the blood vessels and the largest endocrine and metabolic organ of the human body. Its role is to maintain the balance which has the fundamental importance in the life.

Hantavirus pulmonary syndrome real time RT-PCR kit contains a specific ready-to-use system for the detection of Hantavirus pulmonary syndrome by Reverse Transcription Polymerase Chain Reaction (RT-PCR) in the real-time PCR system. The master contains Super Mix for the specific amplification of hanta virus RNA. The reaction is done in one step real time RT-PCR. The first step is a reverse transcription (RT), during which the hanta virus RNA is transcribed into cDNA. Then, a thermostable DNA polymerase is used to amplify the specific gene fragments by polymerase chain reaction. Fluorescence is emitted and measured by the real time systems' optical unit. The detection of amplified Hanta virus (HPS) DNA fragment is performed in fluorimeter channel 530nm with the fluorescent quencher BHQ1. In addition, the kit contains a system to identify possible PCR inhibition by measuring the 560nm fluorescence of the internal control (IC).

#### 4. Kit Contents

Ref.	Type of reagent	Presentation	25rxns
1	HPS Super Mix	1 vial, 350µl	
2	RT-PCR Enzyme Mix	1 vial, 28µl	
3	Molecular Grade Water	1 vial, 400µl	
4	Internal control (IC)	1 vial, 30µl	
5	HPS Positive control(1×10 <sup>7</sup> copies/ml)	1 vial, 30µl	

#### Analysis sensitivity: 1×10<sup>4</sup> copies/ml

**Note:** Analysis sensitivity depends on the sample volume, elution volume, nucleic acid extraction methods and other factors. If you use the RNA extraction kits recommended, the analysis sensitivity is the same as it declares. However, when the sample volume is dozens or even hundreds of times greater than elution volume by some concentrating method, it can be much higher.

#### 5. Storage

- All reagents should be stored at -20°C. Storage at +4°C is not recommended.
- All reagents can be used until the expiration date indicated on the kit label.
- Repeated thawing and freezing (>3x) should be avoided, as this may reduce the sensitivity of the assay.
- Cool all reagents during the working steps.
- Super Mix should be stored in the dark.

#### 6. Additionally Required Materials and Devices

- Biological cabinet
- Real time PCR system
- Desktop microcentrifuge for "eppendorf" type tubes (RCF max. 16,000 x g)
- Vortex mixer
- RNA extraction kit
- Real time PCR reaction tubes/plates
- Cryo-container
- Pipets (0.5 µl – 1000 µl)
- Sterile filter tips for micro pipets
- Sterile microtubes
- Disposable gloves, powderless
- Biohazard waste container
- Refrigerator and Freezer
- Tube racks

#### 7. ⚠ Warnings and Precaution

- Carefully read this instruction before starting the procedure.
- For in vitro diagnostic use only.
- This assay needs to be carried out by skilled personnel.
- Clinical samples should be regarded as potentially infectious materials and should be prepared in a laminar flow hood.
- This assay needs to be run according to Good Laboratory Practice.
- Do not use the kit after its expiration date.
- Avoid repeated thawing and freezing of the reagents, this may reduce the sensitivity of the test.
- Once the reagents have been thawed, vortex and centrifuge briefly the tubes before use.
- Prepare quickly the Reaction mix on ice or in the cooling block.
- Set up two separate working areas: 1) Isolation of the RNA/ DNA and 2) Amplification/ detection of amplification products.
- Pipets, vials and other working materials should not circulate among working units.
- Use always sterile pipette tips with filters.

- Wear separate coats and gloves in each area.
- Do not pipette by mouth. Do not eat, drink, and smoke in laboratory.
- Avoid aerosols

#### 8. Sample Collection, Storage and Transport

- Collected samples in sterile tubes.
- Specimens can be extracted immediately or frozen at -20°C to -80°C.
- Transportation of clinical specimens must comply with local regulations for the transport of etiologic agents.

#### 9. Procedure

##### 9.1 RNA-Extraction

It is necessary to add internal control (IC) in the reaction mix. Internal control (IC) allows the user to determine and control the possibility of PCR inhibition. Add the internal control (IC) 1µl/rxn and the result will be shown in the 560nm Channel.

Nucleic Acid Isolation Kit	Cat. Number	Manufacturer
RNA Isolation Kit	ME-0010/ME-0012	ZJ Biotech
QIAamp Viral RNA Mini Extraction Kit (50)	52904	QIAGEN

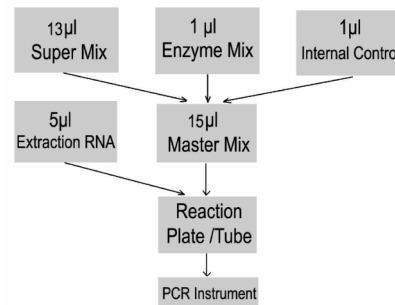
##### 9.2 Internal Control

It is necessary to add internal control (IC) in the reaction mix. Internal control (IC) allows the user to determine and control the possibility of PCR inhibition.

Add the internal control (IC) 1µl/rxn and the result will be shown in the 560nm Channel.

##### 9.3 RT-PCR Protocol

The Master Mix volume for each reaction should be pipetted as follows:



※PCR system without 560nm channel may be treated with 1µl Molecular Grade Water instead of 1µl IC.

- The volumes of Super Mix and Enzyme Mix per reaction multiply with the number of samples, which includes the number of controls and sample prepared. Molecular Grade Water is used as the negative control. For reasons of unprecise pipetting, always add an extra virtual sample. Mix completely then spin down briefly in a centrifuge.
- Pipet 15µl Master Mix with micropipets in sterile filter tips to each real time PCR reaction plate/tubes. Separately add 5µl RNA sample, positive and negative controls to different reaction plate/tubes. Immediately close the plate/tubes to avoid contamination.
- Spin down briefly in order to collect the Master Mix in the bottom of the reaction tubes.
- Perform the following protocol in the instrument:

45°C for 10min	1cycle	Selection of fluorescence channels	
95°C for 15min	1cycle	530nm	Target Nucleic Acid
95°C for 5sec, 60°C for 30sec (Fluorescence measured at 60°C)	40cycles	560nm	IC

**10. Threshold setting:** Choose **Arithmetic** as back ground and **none** as Noise Band method, then adjust the Noise band just above the maximum level of molecular grade water, and adjust the threshold just under the minimum of the positive control.

**11. Quality control:** Negative control, positive control and internal control must be performed correctly, otherwise the sample results is invalid.

Channel	Crossing point value	
	530nm	560nm
Control	Blank	25~35
Molecular Grade Water	Blank	25~35
Positive Control(qualitative assay)	≤35	—

#### 12. Data Analysis and Interpretation

The following sample results are possible:

	Crossing point value		Result Analysis
	530nm	560nm	
1#	Blank	25~35	Below the detection limit or negative
2#	≤38	—	Positive
3#	38~40	25~35	Re-test; If it is still 38~40, report as 1#
4#	Blank	Blank	PCR Inhibition; No diagnosis can be concluded.

For further questions or problems, please contact our technical support at [trade@liferiver.com.cn](mailto:trade@liferiver.com.cn)