## **FIASHTEST**

[Product Name]
Equine Klebsiella (EK) Nucleic Acid Test Kit (Lyophilized)

## cations]

[Intended Use]
The kit uses fluorescent PCR to detect Equine Klebsiella (EK). This product requires the operation of a fluorescent quantitative PCR instrument to achieve rapid detection of POCT.

Testing Principle]
The test kit uses nucleic acid extraction reagents to extract the nucleic acid (DNA/RNA) from the sample. Under the action of 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 action of Taq 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 ST—3' exonuclease activity of Taq polymerase separates the reporting group and quencher group of the fluorescence probe, emitting a specific fluorescence signal. The specific fluorescence signal. The specific fluorescence PCR instrument, and the result is determined based on the Ct value of the sample and the formation of the amplification curve.

Quantity	Storage	
16 pcs	-20°C (Away from light)	
1 pcs		
16 pcs	Room Temperature	
16 pcs		
16 pcs		
	16 pcs 1 pcs 16 pcs 16 pcs	

[Storage conditions and shelf life]
1. Shelf life: 24 months.
2. Production date and expiration date are on the package.

[Compatible Instruments]
This test kit is compatible with FLASHTEST real-time qu
fluorescence PCR instrument.

[Sample] Eye, nose, and throat so wab, anal swab

- [Sample Handling]
  1. Eye, nose, and throat swab: Use a swab to moderately wipe the oral, nasal secretions, or conjunctival secretions.
  2. Anal swab: moisten the swab with dilution solution, then collect the
- 2. Anal swap: nuisten are swap numbers ample.
  3. With the swab in the sample buffer, shake it thoroughly to fully dissolv the pathogen on the swab head into the buffer.

- [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.

[Instructions for Use]

1. Add Elution

1. Add 2DuL of elution from magnetic bead extraction, to each PCR tube Close the lid tightly.

1.2 Shake all the liquid to the bottom of the PCR tube. Use the vortex mixer to mix the PCR tube thoroughly, for 5 seconds. After 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 follow

Step	Temperature	Time	Cycle
1	55°C	3min	1
2	94°C	30s	1
3	94°C 58°C	5s 20s	×40

Channel	FAM	VIC	ROX	Cy5
Target		Internal reference	EK	

# 3. Result Interpretation

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

3.2 Test Result Interpretation			
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 target gene of the virus being tested may lead to false negative results.

Period untervirus being tested may lead to false negative results.

[Product Performance]

1. Positive and negative control consistency: The positive and negative control 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%.

2. Sensitivity: limit 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 55%.

- Sample Is 30 /m.

  [Notes]

  I. Before using a PCR kit, check the lyophilized PCR mix at the bottom of the tube is in good condition (white and clumped). Liquiffied lyophilized PCR mix an 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 mus 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 tips, gloves, and laboratory coats.

  6. After tests, disinfect 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.