

[Product Name] Porcine Circovirus type 2 (PCV2) PCR Test Kit(Lyophilized) [Package Specifica 16 T/box tic s]

[Intended Use] This kill is suitable for the detection of Porcine Circovirus type 2, and the used for the auxiliary diagnosis of clinical Porcine Circovirus type 2 Infection, but it is not for confirmation of the diagnosis. This product requires operation with a fluorescence quantitative PCR instrument at can achieve rapid POCT detection.

can acnieve rapid P/OCI detection. [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 5'--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 using ladetected using a fluorescence PCR instrument, and the result is determined based on the Ct value of the 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

ackage

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

[Sample] EDTA anti-

d n e, thro nt s

[Sample Handling] 1. EDTA anticoagulated blood: Collect blood using a blood collection tube containing EDTA anticoagulant. Add 100 µL of blood to the sample buffer with a disposable dropper. Thoroughly mix the sample buffer with a repetitive pipetting action, using the disposable dropper. 2. Swab: - Nose, throat swab: Use a swab to moderately wipe the oral, nasal coordinor.

secretions. - With the swab in the sample buffer, shake it horoughly to fully dissol the pathogen on the swab head into the buffer. 3. Add 200 µL of mixed buffer to the nucleic acid extraction cartridge fo 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.

Instructions for Use]
1. Add Elution
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1. Add 20L 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
miker 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 follo

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

2.2 Th e is 20

[Channel	FAM	VIC	ROX	Cy5
	Target	Porcine Circovirus type 2	Internal reference		

3. Result Interpretation 3.1 Reference Range:

Parameter	Reference Range	Result Interpretation
Internal	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
	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 shoul 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.

egative results. . Sequence variations caused by mutations or other factors in the target ene of the virus being tested may lead to false negative results.

gene of the virus being tested may lead to false negative results. [Product Performance] 1. Positive and negative control consistency: The positive and negative 1. ontrols 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: Imit of detection is 500 copies/mL. 3. Specificity: This assay does not cross-react with non-target pathogen 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%.

[Notes] 1. Before

[Notes]

 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.
 This product is only for in vitro testing (for animals). All operations mus
strictly follow the instructions.
 Overloading samples may result in false negatives. Retest is
recommended.
 Avoid bubbles in PCR tubes. Keep the tube cap firmly closed.
 Sues dipsoable tips, gloves, and laboratory coats.
 All items in the kit should be treated as biowaste and handled in
accordance with local laboratory regulations.