Probe based qPCR for Gene Detection

Catalog Number Kit Size **TBS42008** 100

DESCRIPTION

The High-Fidelity ASFV TaqProbe qPCR Kit is designed for detecting African swine fever virus DNA from samples in a single PCR reaction using real-time quantitative polymerase chain reaction (qPCR) and probe fluorescence labels. The samples include serum, plasma, fluid, and tissues from swine.

Tribioscience's ASFV Taqprobe qPCR Kit includes a synthesized ASFV DNA fragment as positive controls, negative controls, PCR internal controls from swine derived DNA, internal control probe labeled with Hex, a qPCR super mix, and the primer-probe mix in which the probe has been labeled with FAM for the target gene. These aid in a straightforward interpretation of the results.

KEY FEATURES

High sensitivity and specificity for target detection.

- High efficiency: the optimal systemic conditions for PCR amplification.
- Streamlined protocol: Just add DNA Template and water.
- ♦ No cross reactivity with others.

KIT CONTENTS

Name	Unit Size
HF qPCR Enymix P1	0.8mL
HF qPCR Enymix P2	0.5mL
HF qPCR Enymix P3	60µL
HF qPCR Enymix P4	60µL
HF qPCR Enymix P5	0.5mL

STORAGE CONDITION

The kit is shipped on ice and stored at -20°C for long-term storage. Shelf life of 12 months after receipt.

PCR PROTOCOL

1	. Set up PCI	R reaction	for each	sample	in 20	μL
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Reaction Component	Volume (µL)
HF qPCR Enymix P1	7.0
HF qPCR Enymix P2	4.0
HF qPCR Enymix P5	4.0
DNA sample	5.0
Final Volume	20 µL

Internal control should be included as below: HF qPCR Enymix P3 or P4 (4µL/reaction).

The ASFV probe was labeled with Fam, and Internal control probe was labeled with HEX.

2. Suggested PCR conditions

	Amplification	PCR		
Step	HOLD	CYCLE (40x cycles)		
		Denature	Anneal/ Extend	
Temperature	95°C	95°C	60°C	
Time	1 min	15 sec	60 sec	

DATA ANALYSIS

Positive Reaction: Sample Ct \leq 37 w/ Positive, Negative and Blank controls normal.

Negative Reaction: Sample Ct \ge 38 w/ Positive, Negative and Blank controls normal.

PCR internal control is positive in all samples, positive and negative controls. The positive response indicates a normal PCR amplification. Otherwise, the PCR reaction may be inhibited.

Repeat Reaction: If one of the control reactions is not normal, PCR reaction is failed, and should be repeated.

For research use only.