Ruminant

# **VDRG® FMDV 3Diff/PAN Ag** Rapid Kit

Asia1

SAT1

SAT2

SAT2

С



Foot and Mouth Disease Virus	pigs and various wildlife to high mutation rates v a vesicular condition of	e species. There are sev which constantly gener the feet, buccal mucos stic reagent can detect	cloven hoofed (two-toed ven types (O, A, C, SAT 1, ate new FMDV variants. T sa and, in females, the ma FMDV specific serotype a pping the specimens.	SAT 2, SAT 3 and Asia ypical cases of FMD a immary glands.	a1), that are subject re characterized by	
ntroduction	- Spec	All serotype antigens cific serotypes : O, A, A erotypes : O, A, C, Asia rromatographic assay fluid, Infected tissue, s g Rapid Test Device	detection Asia1 a1, SAT1, SAT2, and SAT3		Contraction of the second seco	
Features	<ul> <li>Differential diagnosis of 3 different serotypes (O,A,Asia1) of FMD virus</li> <li>Available Rapid kit at field farm condition</li> <li>Applicable to various suspect specimens (vesicular fluid, infected tissue, saliva, and cultivated virus etc)</li> <li>Concurrent diagnosis both of a virus's common antigen(all 7 serotypes) and serotype-specific antigens (O,A,Asia1)</li> <li>Clinical Sensitivity <ol> <li>FMD type 0 88.2%(n=60/68) vs PCR</li> <li>FMD type A 100%(n=70), Asia1 100%(n=62), SAT1 100%(n=20), SAT2 100%(n=20), SAT3 100%(n=20), C 100%(n=20), in spiking samples with virus (titer : 1.0x10<sup>5</sup>TCID<sub>50</sub>/mL or higher)</li> <li>Clinical Specificity <ol> <li>type O, type A and strip PAN : normal cattle 100%(n=92), normal pigs 100%(n=400)</li> <li>type Asia1 : normal cattle 100%(n=92), normal pigs 99.5%(n=398/400)</li> </ol> </li> </ol></li></ul>					
Cross reactivity	VDRG <sup>®</sup> FMDV 3Diff/PAN Ag Rapid kit					
Cross reactivity among	Serotype	0	3Diff A	Asia1	PAN	
serotypes of FMDV	0	Positive	Negative	Negative	Positive	
	A	Negative	Positive	Negative	Positive	

Negative

Negative

Negative

Negative

Negative

Positive

Negative

Negative

Negative

Negative

Negative

Negative

Negative

Negative

Negative

Positive

Positive

Positive

Positive

Positive

# Sample preparation

### A. Sample collection

- 1. Collection of fluid from intact vesicle : draw vesicular fluid with syringe.
- 2. Collection of fluid from ruptured vesicles : Soak vesicular fluid using a cotton swab.
- 3. Tissue sampling from ruptured lesions :
- ① Follow the instruction manual in VDRG® Tissue Sample Extraction kit (CAT.NO. EXT-TIS-11, not provided).

② Collect 0.2g of fresh and friable epithelium (nail size of little finger) from surface or margin of vesicles or other tissues of interest.

- 4. Saliva : Collect saliva from swine or bovine using appropriate method.
  - ① For bovine, collect saliva directly from tongue using disposable plastic gloves.
- (2) For swine, collect saliva using chewing rope or other oral fluid collection kit.
- 5. Cultured virus : Collect virus culture media using micropipette.

### B. Sample processing

- 1. Syringe-collected fresh vesicular fluid
- 1 Add 1 scale (approximately 250ul) of Sample Dilution Buffer to the test tube using dropper.
- 2 Add 250 $\mu\ell$  of syringe-collected vesicular fluid to the test tube and mix gently.
- 2. Swab-collected fluid from ruptured vesicles
  - 1 Add 2 scales (approximately 500ul) of Sample Dilution Buffer to the test tube using dropper.
  - (2) Soak the swab in the dilution buffer, mix by swirling and extract the vesicular lesion fluid by pressing the cotton swab against the tube wall.
  - ③ Remove the swab from the test tube after extraction.
- 3. Tissue-extracted fluid
  - ① Follow the instruction manual in VDRG® Tissue Sample Extraction Kit (Cat.No. EXT-TIS-11, not provided).
  - 2 Add 4 scales (1mL) of Sample Dilution Buffer to the extraction vial.
  - 3 Add tissue sample to the extraction vial.
  - 4 Cut the tissue into pieces using scissors and grind using pestle and sand included in the kit.
  - (5) Leave the homogenate for 2~3 minutes to settle down tissue fraction.
  - <sup>(6)</sup> Use the clarified fluid for testing.
- 4. Saliva

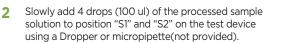
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- 1 Add 2 scales (approximately 500ul) of Sample Dilution Buffer to the test tube using the dropper.
- (2) Centrifugate (6,000rpm, 10min) the collected saliva and soak the swab with supernatant of centrifugated saliva.
- (3) Soak the swab in the dilution buffer, mix by swirling and extract the saliva by pressing the cotton swab against the tube wall.
- ④ Remove the swab from the test tube after extraction.
- (5) Use the diluted and clarified saliva for testing.
- 5. Virus culture media
- 1 Add 200ul of Sample Dilution Buffer to the test tube, Eppendorf tube or microplates.
- ② Add 200ul of virus culture media to the test tube and mix by several times of swirling.

## Test procedure

Add 1-4 scales of Sample Dilution Buffer depending on sample types to Test Tube using Dropper and process the sample.







**3** Read results within **15 minutes** exactly. Reading later than 15 minutes may cause inaccurate results.

1. Positive when both control and test lines are red.



1) FMDV serotype O positive result



 FMDV serotype Asia-1 positive result



2) FMDV serotype A positive result



4) FMDV other serotype (SAT1, SAT2, SAT3, C) positive result



3. Re-test when control line is not visible.

2. Negative when only control lines are red.



#### **Order Information**

Cat No.	Product Name	Quantity	
PM-FMD-16	VDRG <sup>®</sup> FMDV 3Diff/PAN Ag Rapid kit	10 Tests/Box	