

For research use only

Cat. No. INT-IP11527 | 48 Tests

Porcine Circovirus Type 2 Detection Kit

Test for the detection of Porcine Circovirus type-2 by one-step
PCR

User Manual

REV.2.2



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Table of Contents

1.	Description	1
2.	Storage	1
3.	Contents	1
4.	Specimen	1
5.	Additional required materials	2
6.	Procedure	2
6.1	DNA preparation.....	2
6.2	Amplification	2
6.3	Detection of amplification product	3
6.4	Interpretation.....	3
6.5	Elimination of carry-over contamination	3
7.	Notice	4
8.	Trouble shooting	4
9.	Ordering information	4

1. DESCRIPTION

Postweaning multisystemic wasting syndrome (PMWS) was initially recognized in weaning piglets of a high-health-status herd. Since then, PMWS has become an economically important disease in virtually all regions of the world that produce pigs. PMWS primarily affects pigs between 5 and 18 weeks of age.

Clinical PMWS signs include progressive weight loss, dyspnea, tachypnea, anemia, diarrhea, and jaundice. The primary causative agent of PMWS has been identified as porcine circovirus type 2 (PCV2). Porcine circovirus type 1 (PCV1) was discovered as a noncytopathic contaminant of the porcine kidney cell line PK-15 and is nonpathogenic to pigs.

Both PCV1 and PCV2 are small, nonenveloped viruses with a single-stranded circular DNA genome of about 1.76 kb. PCV belongs to the *Circoviridae* family along with chicken anemia virus, psittacine beak and feather disease virus, and tentative members columbid circovirus, goose circovirus, and canary circovirus.

Porcine Circovirus type 2 Detection Kit is direct detection of porcine circovirus(type-1 and type-2) on the basis of a genetic database, so it can diagnose very fast and accurately. It can amplify only specific gene using the PCR (Polymerase Chain Reaction) method, and take only 2~3 hours for detection. Therefore, it is a very fast, accurate and reliable technique.

2. STORAGE

The components of **Porcine Circovirus type 2 Detection Kit** should be stored at -20 °C, under this condition, the kit is stable until expiration date stated on the label.

3. CONTENTS

Porcine Circovirus type 2 PCR Pre-mixture	48 tubes
DNase/RNase-free water (white cap)	1 vial
PCV2 positive control (Yellow cap)	2 vial

Component in 20 µl reaction

i-StarTaq™ DNA Polymerase
dNTPs
PCR Reaction buffer
Chemical stabilizer
Gel loading buffer
8-MOP (dissolved in DMSO)
Primers for PCV type-2

4. SPECIMEN

Performs the test with whole blood, lymphoid tissues, lung, tonsil, nasal or tonsil swab. The specimen should be stored at -20 °C prior to use.

5. ADDITIONAL REQUIRED MATERIALS

- Disposable gloves
- DNA extraction kit (see 6.1 DNA preparation method)
- Pipettes
- Sterile pipette tip
- Vortex mixer
- Centrifuge for microcentrifuge tubes
- Thermal cycler
- Electrophoresis kit
- UV transilluminator

6. PROCEDURE

Please read through the entire procedure before starting.

6.1 DNA Preparation

Various manufacturers offer DNA isolation kits. Please carry out the DNA isolation according to the manufacturer's instructions. The following standard extraction kit is recommended.

Product	Catalog No.	Manufacturer
Viral gene-spin™ Viral DNA/RNA Extraction Kit	17151	iNtRON Biotechnology, Inc.

6.2 Amplification

- ① Prepare appropriate PCR premix tubes and label. And one PCR premix tube for positive control.
- ② Add 2 μl of template DNA into the PCR premix tube.
- ③ Add 18 μl of DNase/RNase-free water into the PCR premix tube to total volume as 20 μl.
- ④ Add 2 μl of positive control and 18 μl of RNase-free water into a PCR premix tube for monitoring of amplification and easy interpretation.
- ⑤ Dissolve the blue pellet by pipetting.
Note :The pellet is easily dissolved, by letting the mixture stand at R.T. for 1-2minutes after adding water.
- ⑥ (Optional) Add mineral oil. This step is unnecessary when using a thermal cycler that employs a top heating method (general methods).
- ⑦ Perform PCR reaction of samples as the below process using PCR machine.

PCR cycle		Temp.	Time
1 Cycle	Initial Denaturation	94 °C	5 min.
	Denaturation	94 °C	30 sec.
40 Cycles	Annealing	54 °C	30 sec.
	Extension	72 °C	40 sec.
1 Cycle	Final extension	72 °C	5 min.

6.3 Detection of Amplified Products

- ① Prepare 1.5% agarose gel containing RedSafe™ Nucleic Acid Staining Solution (20,000x).
- ② Load 7 μl of PCR product and positive control on agarose gel without adding a loading-dye buffer and perform electrophoresis.
- ③ Run electrophoresis by 100V (required about 30~40 minutes).
- ④ Identify the result on ultra-violet (UV) transilluminator.

6.4 Interpretation

- Expected PCR product size : 492 bp

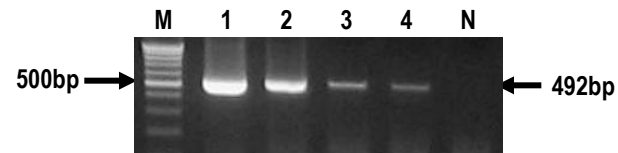


Fig 1. Electrophoresis of PCR product by **Porcine Circovirus type 2 Detection Kit**
 Lane M : 100bp Molecular ladder (iNtRON Biotechnology)
 Lane 1~4 : PCV type-2 positive sample
 Lane N : Negative control

6.5 Elimination of carry-over contamination

- Each PCR/RT-PCR Pre-mixture contains 8-methoxypsoralen (8-MOP) for preventing of carry-over contamination.
- All PCR products should be discarded after UV irradiation (10 min/365nm) for preventing from carry-over contamination.

