

## 1. DESCRIPTION

Psitticine Beak and Feather Disease (PBFVD) is a chronic disease characterized by feather dystrophy and loss, beak deformity and ultimately, death. It is caused by a non-enveloped icosahedral DNA virus (PBFVDV) belonging to the family Circoviridae. The disease has been reported in Australia, North America, Europe and Asia. PBFVD virus usually infects birds less than 3 years of age. The virus is spread from mother to egg or directly to chicks. Viral particles can be spread in feather dust carried by air currents, dried feces or even on the clothing of human handlers. Nest materials, feeding formula, feeding utensils, nets, bird carriers, food dishes and other fomites are easily contaminated with this virus. The first clinically detectable sign of PBFVD is the appearance of necrotic, abnormally formed feathers. Many birds infected with PBFVD die within 6-12 months of onset of clinical signs. However, some birds have been known to survive 10-15 years and become chronic carriers. Death usually occurs from secondary bacterial, fungal, parasitic, chlamydial, or viral infections.

VetPCR™ PBFVDV Detection Kit is the direct detection of Psitticine Beak and Feather Disease 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 3 hours for detection. Therefore, it is a very fast, accurate and reliable technique.

## 2. STORAGE

The components of VetPCR™ PBFVDV Detection Kit should be stored at -20°C. Under this condition, the kit is stable until expiration date stated on the label.

## 3. CONTENTS

	Kit 48	Kit 96	
VetPCR™ PBFVDV PCR Pre-mixture .....	48	96	tubes
DNase/RNase-free water .....	1	1	vial
PBFVDV PCR Positive control .....	1	1	vial
PBFVDV PCR Positive control Pre-mixture .....	4	8	tubes
Brig™ Molecular Weight marker .....	1	1	vial
Mineral Oil .....	1	1	vial
DNA purification kit (see step 6.1) .....	50	100	tests

## 4. SPECIMEN

0.5 ml whole blood in EDTA (purple top) tube, or cloacal swab, or swab of the outer surface of liver, spleen or kidney, or tissue.

## 5. ADDITIONAL REQUIRED MATERIALS

- Disposable gloves
- 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 manufacturers instructions. The following standard DNA Purification kit is recommended.

Product	Catalog No.	Manufacturer
Bioingentech™ Genomic DNA Purification Kit (50 test)	230040(50)	Bioingentech Biotechnology Inc.
Bioingentech™ Genomic DNA Purification Kit (100 test)	230040(100)	Bioingentech Biotechnology Inc.

### 6.2 AMPLIFICATION

1.- Prepare appropriate PCR Premix tubes and one PCR Premix tube for Positive control. Label.

2.- Add 6µl of DNase/RNase-free water into the PCR Premix tube to total volume as 11µl.

3.- Add 2µl of template DNA into the PCR Premix tube to total volume as 13µl.

4.- Add 6µl of DNase/RNase-free water and 2µl of PCR Positive control into a PCR Positive control Premix tube for monitoring of amplification and easy interpretation.

5.- Add mineral oil (11µl). This step is necessary, even when using a thermal cycler that employs a top heating method.

6.- Perform PCR reaction of samples as the below process, using a PCR thermal cycler.

PCR cycle		Temp.	Time
30 Cycles	Initial Denaturation	94°C	2 min.
	Denaturation	94°C	30 sec.
	Annealing	55°C	30 sec.
1 Cycle	Extension	72°C	30 sec.
	Final extension	72°C	5 min.

### 6.3 DETECTION OF AMPLIFIED PRODUCTS

1.- Prepare 1.5% agarose gel containing Ethidium bromide (Et-Br).

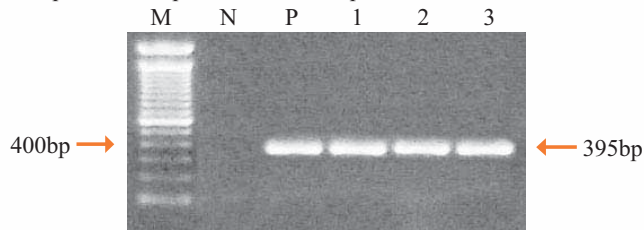
2.- Load 7µl of PCR product, 7µl of positive control and 2µl of Brig™ Molecular Weight marker on agarose gel without adding a loading-dye buffer and perform electrophoresis.

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 :395bp



**Fig1.** Electrophoresis of PCR product by VetPCR™ PBFVDV Detection Kit  
Lane M : Brig™ Molecular Weight Marker (Bioingentech Ltd.)  
Lane N : Negative control  
Lane P : Positive control  
Lane 1~3 : PBFVDV Positive sample

## 7. NOTICE

- For research purpose only. Not for use in diagnostic procedures for clinical purposes. *For in Vitro Use Only.*
- Take care in handling of specimen to minimize risk of infection.
- The PCR process is covered by patents issued and applicable in certain countries. Bioingentech Biotechnology Inc. does not encourage or support the unauthorized or unlicensed use of the PCR process. Use of this product is recommended for persons that either have a license to perform PCR or are not required to obtain a license.

## 8. TROUBLE SHOOTING

- 1.- In the case of difficult to interpret results due to non-specific bands; reduce amount of template by 1/10 dilution, heated at 65° C for 5 min. and reacts again.
- 2.- Preparation of PCR reaction at room temperature may cause the non-specific band.
- 3.- All procedure should be carried out on ice.

## 9. ORDERING INFORMATION

Product	Catalog No.
VetPCR™ PBFVDV Detection Kit 48	VET0014A(48)
VetPCR™ PBFVDV Detection Kit 96	VET0014A(96)
Brig™ Molecular Weight Marker	24012



**Bioingentech Ltd.**

Salas 350, piso 2, Concepción, Chile  
Telephone (56)-(41)-2790435  
Fax (56)-(41)-2790435  
info@bioingentech.com  
www.bioingentech.com

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