# Aftovaxpur DOE (80) O Taiwan 3/97 + Asia1 Shamir + SAT2 Saudi Arabia

Not authorised

- Foot-and-mouth disease virus, serotype O, strain Taiwan 3/97, Inactivated
- Foot-and-mouth disease virus, serotype Asia 1, strain Shamir, Inactivated
- Foot-and-mouth disease virus, serotype SAT2, strain Saudi Arabia, Inactivated

# Product identification

#### Medicine name:

Aftovaxpur DOE (80) O Taiwan 3/97 + Asia1 Shamir + SAT2 Saudi Arabia

#### **Active substance:**

Foot-and-mouth disease virus, serotype O, strain Taiwan 3/97, Inactivated Foot-and-mouth disease virus, serotype Asia 1, strain Shamir, Inactivated Foot-and-mouth disease virus, serotype SAT2, strain Saudi Arabia, Inactivated

### **Target species:**

This information is not available for this product.

#### Route of administration:

Intramuscular use Subcutaneous use

### **Product details**

### **Active substance and strength:**

Foot-and-mouth disease virus, serotype O, strain Taiwan 3/97, Inactivated

Presentation strength:? 6 PD50 Reference:Hse Index:0

Foot-and-mouth disease virus, serotype Asia 1, strain Shamir, Inactivated

Presentation strength:? 6 PD50 Index:11

Foot-and-mouth disease virus, serotype SAT2, strain Saudi Arabia, Inactivated

Presentation strength:? 6 PD50 Index:12

#### **Pharmaceutical form:**

Emulsion for injection

### Anatomical therapeutic chemical veterinary (ATCvet) codes:

**Q102AA04** 

### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Surrendered

#### Authorised in:

Austria, Belgium, Bulgaria, Croatia, Cyprus, Czechia, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, United Kingdom (Northern Ireland)

# Additional information

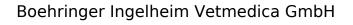
# **Entitlement type:**

Marketing Authorisation

# Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

# Marketing authorisation holder:



### Marketing authorisation date:

15/07/2013

### Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France Boehringer Ingelheim Animal Health France

## **Responsible authority:**

**European Commission** 

#### **Authorisation number:**

This information is not available for this product.

## Date of authorisation status change:

16/05/2023

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

# **Documents**

Combined File of all Documents

English (PDF)

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