# Aftovaxpur DOE (51) O Taiwan 3/97 + A22 Iraq + A24 Cruzeiro

Not authorised

- Foot-and-mouth disease virus, serotype O, strain Taiwan 3/97, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A22
  Iraq, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A24 Cruzeiro, Inactivated

# **Product identification**

#### **Medicine name:**

Aftovaxpur DOE (51) O Taiwan 3/97 + A22 Iraq + A24 Cruzeiro

#### **Active substance:**

Foot-and-mouth disease virus, serotype O, strain Taiwan 3/97, Inactivated Foot-and-mouth disease virus, serotype A, strain A22 Iraq, Inactivated Foot-and-mouth disease virus, serotype A, strain A24 Cruzeiro, 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 A, strain A22 Iraq, Inactivated

Presentation strength:? 6 PD50 Index:11

Foot-and-mouth disease virus, serotype A, strain A24 Cruzeiro, 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

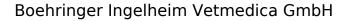
# **Entitlement type:**

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)

Published on: 19/03/2024

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