

# Aftovaxpur DOE (29) O1 Manisa + O1 BFS + A22 Iraq

Not  
authorised

- Foot-and-mouth disease virus, serotype O, strain O1 Manisa, Inactivated
- Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A22 Iraq, Inactivated

## Product identification

### **Medicine name:**

Aftovaxpur DOE (29) O1 Manisa + O1 BFS + A22 Iraq

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### **Active substance:**

Foot-and-mouth disease virus, serotype O, strain O1 Manisa, Inactivated

Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated

Foot-and-mouth disease virus, serotype A, strain A22 Iraq, Inactivated

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### **Target species:**

This information is not available for this product.

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### **Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

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

Foot-and-mouth disease virus, serotype O, strain O1 Manisa, Inactivated

Presentation\_strength:? 6 PD50 Reference:Hse Index:0

Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated

Presentation\_strength:? 6 PD50 Index:11

Foot-and-mouth disease virus, serotype A, strain A22 Iraq, Inactivated

Presentation\_strength:? 6 PD50 Index:12

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### **Pharmaceutical form:**

Emulsion for injection

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

QI02AA04

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### **Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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### **Authorisation status:**

Surrendered

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### **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)

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## Additional information

### **Entitlement type:**

Marketing Authorisation

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### **Legal basis of product authorisation:**

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

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### **Marketing authorisation holder:**

Boehringer Ingelheim Vetmedica GmbH

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**Marketing authorisation date:**

15/07/2013

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France

Boehringer Ingelheim Animal Health France

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

16/05/2023

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To consult adverse reactions on veterinary medicinal products please go to

[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

Published on: 19/03/2024

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