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

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

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 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

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

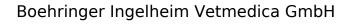
Entitlement type:

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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