

Aftovaxpur DOE (39) O1 Manisa + A24 Cruzeiro + A Turkey 14/98

Not
authorised

- Foot-and-mouth disease virus, serotype O, strain O1 Manisa, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A24 Cruzeiro, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A/Turkey/14/98, Inactivated

Product identification

Medicine name:

Aftovaxpur DOE (39) O1 Manisa + A24 Cruzeiro + A Turkey 14/98

Active substance:

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

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

Foot-and-mouth disease virus, serotype A, strain A/Turkey/14/98, 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 A, strain A24 Cruzeiro, Inactivated

Presentation_strength:? 6 PD50 Index:11

Foot-and-mouth disease virus, serotype A, strain A/Turkey/14/98, Inactivated

Presentation_strength:? 6 PD50 Index:12

Pharmaceutical form:

Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AA04

Legal status of supply:

Medicinal product subject to medical 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:

Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:

15/07/2013

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France
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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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