Aftovaxpur DOE (41) O1 Manisa + A Turkey 14/98 + Asia1 Shamir

Not authorised

- Foot-and-mouth disease virus, serotype O, strain O1 Manisa, Inactivated
- Foot-and-mouth disease virus, serotype A, strain Turkey 14/98, Inactivated
- Foot-and-mouth disease virus, serotype Asia 1, strain Shamir, Inactivated

Product identification

Medicine name:
Aftovaxpur DOE (41) O1 Manisa + A Turkey 14/98 + Asia1 Shamir

Active substance:
Foot-and-mouth disease virus, serotype O, strain O1 Manisa, Inactivated
Foot-and-mouth disease virus, serotype A, strain Turkey 14/98, Inactivated
Foot-and-mouth disease virus, serotype Asia 1, strain Shamir, 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 Turkey 14/98, Inactivated

Presentation_strength:? 6 PD50 Index:11

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

Presentation_strength:? 6 PD50 Index:12

Pharmaceutical form:

Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AA04

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:

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

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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ema-puar-aftovaxpur-v-2292-var-ii-0001-en.pdf

ema-puar-aftovaxpur-v-2292-par-en.pdf

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