Aftovaxpur DOE (36) O1 Manisa + O Taiwan 3/97 + Asia1 Shamir

Not authorised

- Foot-and-mouth disease virus, serotype O, strain O1 Manisa, Inactivated
- Foot-and-mouth disease virus, serotype O, strain Taiwan 3/97, Inactivated
- Foot-and-mouth disease virus, serotype Asia 1, strain Shamir, Inactivated

Product identification

Medicine name:
Aftovaxpur DOE (36) O1 Manisa + O Taiwan 3/97 + Asia1 Shamir

Active substance:
Foot-and-mouth disease virus, serotype O, strain O1 Manisa, Inactivated
Foot-and-mouth disease virus, serotype O, strain Taiwan 3/97, 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 O, strain Taiwan 3/97, 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
Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France
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
Download

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

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

ema-puar-aftovaxpur-v-2292-var-ii-0009-en.pdf