Aftovaxpur DOE (43) O1 BFS + O Taiwan 3/97 + A24 Cruzeiro

Authorised

- Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated
- Foot-and-mouth disease virus, serotype O, strain Taiwan 3/97, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A24 Cruzeiro, Inactivated

Product identification

Medicine name:

Aftovaxpur DOE (43) O1 BFS + O Taiwan 3/97 + A24 Cruzeiro

Active substance:

Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated Foot-and-mouth disease virus, serotype O, strain Taiwan 3/97, Inactivated Foot-and-mouth disease virus, serotype A, strain A24 Cruzeiro, Inactivated

Target species:

Cattle

Sheep

Pig

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Foot-and-mouth disease virus, serotype O, strain O1 BFS, 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 A, strain A24 Cruzeiro, Inactivated

Presentation strength:≥ 6 PD50 Index:12

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- . Cattle
 - Not applicable. 0 day Zero days
- Sheep
 - Not applicable. 0 day Zero days
- Pig
 - Not applicable. 0 day Zero days

Subcutaneous use:

- . Cattle
 - Not applicable. 0 day Zero days
- Sheep
 - Not applicable. 0 day Zero days
- Pig
 - Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

Q102AA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

This information is not available for this product.

Package description:

Packaging:Bottle (polypropylene), Package_size:1 bottle, Content:300 ml Packaging:Bottle (polypropylene), Package_size:1 bottle, Content:20 ml Packaging:Bottle (polypropylene), Package_size:1 bottle, Content:50 ml Packaging:Bottle (polypropylene), Package_size:10 bottles, Content:100 ml Packaging:Bottle (polypropylene), Package_size:10 bottles, Content:300 ml Packaging:Bottle (polypropylene), Package_size:10 bottles, Content:50 ml Packaging:Bottle (polypropylene), Package_size:10 bottles, Content:200 ml Packaging:Bottle (polypropylene), Package_size:10 bottles, Content:200 ml Packaging:Bottle (polypropylene), Package_size:10 bottles, Content:200 ml Packaging:Bottle (polypropylene), Package_size:1 bottle, Content:20 ml Packaging:Bottle (polypropylene), Package_size:1 bottle, Content:100 ml

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

European Commission
Authorisation number: This information is not available for this product.
Date of authorisation status change: 15/07/2013
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-0009-en.pdf
ema-puar-aftovaxpur-v-2292-par-en.pdf
ema-puar-aftovaxpur-v-2292-var-ii-0001-en.pdf

Responsible authority:

Source URL: https://medicines.health.europa.eu/veterinary/60000000939