

Aftovaxpur DOE (70) O1 Manisa + A Turkey 14/98 + SAT2 Saudi Arabia

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 SAT2, strain Saudi Arabia, Inactivated

Product identification

Medicine name:

Aftovaxpur DOE (70) O1 Manisa + A Turkey 14/98 + SAT2 Saudi Arabia

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 SAT2, strain Saudi Arabia, 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 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 SAT2, strain Saudi Arabia, 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:

QI02AA04

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

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

7/04/2014

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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