File downloaded on 2026-11-08
Source URL: https://medicines.health.europa.eu/veterinary/en/600000000322

Aftovaxpur DOE (WD) (20) O Taiwan 3/97 + A24 Cruzeiro

Not authorised

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

Product identification

Medicine name:
Aftovaxpur DOE (WD) (20) O Taiwan 3/97 + A24 Cruzeiro

Active substance:
Foot-and-mouth disease virus, serotype A, strain O/Taiwan/3/97, Inactivated
Foot-and-mouth disease virus, serotype A, strain A24 Cruzeiro, Inactivated
Foot-and-mouth disease virus, serotype A, strain O/Taiwan/3/97, Inactivated
Foot-and-mouth disease virus, serotype A, strain A24 Cruzeiro, Inactivated

Target species:
Cattle
Sheep
Pig
Cattle

Sheep
Pig

Route of administration:
Intramuscular use
Subcutaneous use
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Foot-and-mouth disease virus, serotype A, strain O/Taiwan/3/97, 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 O/Taiwan/3/97, 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

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

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:
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
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
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
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
Boehringer Ingelheim Animal Health France

Responsible authority:
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: 14/03/2026
Download

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

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

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