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Aftovaxpur DOE (WD) (18) O1 BFS + Asia1 Shamir

Non autorizzato

- Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated
- Foot-and-mouth disease virus, serotype Asia 1, strain Shamir, Inactivated
- Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated
- Foot-and-mouth disease virus, serotype Asia 1, strain Shamir, Inactivated

Identificazione del prodotto

Denominazione del medicinale:
Aftovaxpur DOE (WD) (18) O1 BFS + Asia1 Shamir

Sotsanza attiva:
Disponibile solo in English
Disponibile solo in English
Disponibile solo in English
Disponibile solo in English

Specie di destinazione:
bovini
Ovino
Suino
bovini
Ovino

Suino

Via di somministrazione:
Uso intramuscolare
Uso sottocutaneo
Uso intramuscolare
Uso sottocutaneo

Informazioni sul prodotto

Sostanza attiva e dosaggio:
Disponibile solo in English
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Disponibile solo in English
Presentation_strength:≥ 6 PD50 Index:11
Disponibile solo in English
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Disponibile solo in English
Presentation_strength:≥ 6 PD50 Index:11

Forma farmaceutica:
Emuslione iniettabile

Tempo di attesa per via di somministrazione:
Uso intramuscolare:

- **bovini**
  - non applicabile. 0 giorno Zero days
- **Ovino**
  - non applicabile. 0 giorno Zero days
- **Suino**

- non applicabile. 0 giorno Zero days

Uso sottocutaneo:

- **bovini**
  - non applicabile. 0 giorno Zero days
- **Ovino**
  - non applicabile. 0 giorno Zero days
- **Suino**
  - non applicabile. 0 giorno Zero days

Uso intramuscolare:

- **bovini**
  - non applicabile. 0 giorno Zero days
- **Ovino**
  - non applicabile. 0 giorno Zero days
- **Suino**
  - non applicabile. 0 giorno Zero days

Uso sottocutaneo:

- **bovini**
  - non applicabile. 0 giorno Zero days
-

Ovino

- non applicabile. 0 giorno Zero days

•

Suino

- non applicabile. 0 giorno Zero days

Codice anatomico, terapeutico e chimico veterinario (codice ATCvet):
QI02AA04

Stato legale della fornitura:
Medicinale veterinario soggetto a prescrizione veterinaria

Stato dell’autorizzazione:
Rinunciato

Autorizzato in:
Austria , Belgio , Bulgaria , Croazia , Cipro , Repubblica Ceca , Danimarca , Estonia , Finlandia , Francia , Germania , Grecia , Ungheria , Islanda , Irlanda , Italia , Lettonia , Liechtenstein , Lituania , Lussemburgo , Malta , Paesi Bassi , Norvegia , Polonia , Portogallo , Romania , Slovacchia , Slovenia , Spagna , Svezia ,

Disponibile solo in Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Ulteriori informazioni

Tipo di diritto:

Base giuridica dell’autorizzazione del prodotto:
Applicazione completa (articolo 12,paragrafo 3 ,della Direttiva N°2001/82/CE)

Titolare dell’autorizzazione all’immissione in commercio:
Boehringer Ingelheim Vetmedica GmbH

Data di autorizzazione all'immissione in commercio:
15/07/2013

Siti di fabbricazione e rilascio dei lotti:

Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France
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Autorità responsabile:
European Commission

Numero di autorizzazione:
Queste informazioni non sono disponibili per questo prodotto.

Data della modifica dello stato dell’autorizzazione:
15/07/2013

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documenti

File combinato di tutti i documenti

italiano (PDF)
Pubblicato il: 14/03/2026
Scaricamento

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

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