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Aftovaxpur DOE (WD) (02) O1 BFS

Nav atļautas

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

Zāļu identifikācija

Zāļu nosaukums:
Aftovaxpur DOE (WD) (02) O1 BFS

Aktīvā viela:
Pieejamas tikai English
Pieejamas tikai English

Mērķsugas:
Liellops
Aita
Cūka
Aita
Cūka
Liellops

Lietošanas veids:
intramuskulārai lietošanai
subkutānai lietošanai
subkutānai lietošanai

intramuskulārai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:
Pieejamas tikai English
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Pieejamas tikai English
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0

Farmaceitiskā forma:
Emulsija injekcijām

Ierobežojumu periods pēc lietošanas veida:
intramuskulārai lietošanai:

- **Liellops**
  - Nav piemērojams. 0 diena Zero days
- **Aita**
  - Nav piemērojams. 0 diena Zero days
- **Cūka**
  - Nav piemērojams. 0 diena Zero days

subkutānai lietošanai:

- **Liellops**
  - Nav piemērojams. 0 diena Zero days
- **Aita**

- Nav piemērojams. 0 diena Zero days

- **Cūka**
  - Nav piemērojams. 0 diena Zero days

subkutānai lietošanai:

- **Aita**
  - Nav piemērojams. 0 diena Zero days

- **Cūka**
  - Nav piemērojams. 0 diena Zero days

- **Liellops**
  - Nav piemērojams. 0 diena Zero days

intramuskulārai lietošanai:

- **Liellops**
  - Nav piemērojams. 0 diena Zero days

- **Aita**
  - Nav piemērojams. 0 diena Zero days

- **Cūka**
  - Nav piemērojams. 0 diena Zero days

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QI02AA04

Izplatīšanas juridiskie nosacījumi:

Recepšu veterinārās zāles

Atļaujas statuss:

Atsaukta

Atļautas:

Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:

Pieejamas tikai English Italian

Tirdzniecības atļaujas turētājs:

Boehringer Ingelheim Vetmedica GmbH

Tirdzniecības atļaujas datums:

15/07/2013

Ražošanas vietas sēriju izlaidei:

Boehringer Ingelheim Animal Health France
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Atbildīgā iestāde:
European Commission

Atļaujas numurs:
Šī informācija par šīm zālēm nav pieejama.

Atļaujas statusa maiņas datums:
15/07/2013

Lai uzzinātu par veterināro zāļu blakusparādībām, lūdzu, dodieties uz: www.adrreports.eu/vet

Dokumenti

Visu dokumentu apvienotais fails

latviešu (PDF)
Publicēts: 14/03/2026
Lejupielādēt

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

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

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