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Aftovaxpur DOE (WD) (07) Asia1 Shamir

Ei myönnetty

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

Valmisteen perustiedot

Lääkkeen nimi:
Aftovaxpur DOE (WD) (07) Asia1 Shamir

Vaikuttava aine:
Saatavissa vain kielillä English
Saatavissa vain kielillä English

Kohde-eläinlajit:
Nauta
Lammas
Sika
Nauta
Lammas
Sika

Antoreitti:
Lihakseen
Ihon alle
Ihon alle

Lihakseen

Valmistetiedot

Vaikuttava aine ja vahvuus:
Saatavissa vain kielillä English
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Saatavissa vain kielillä English
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0

Lääkemuoto:
Injektioneste, emulsio

Varoaika antoreiteittäin:
Lihakseen:

- **Nauta**
  - Not applicable. 0 day Zero days
- **Lammas**
  - Not applicable. 0 day Zero days
- **Sika**
  - Not applicable. 0 day Zero days

Ihon alle:

- **Nauta**
  - Not applicable. 0 day Zero days
- **Lammas**

- Not applicable. 0 day Zero days

- **Sika**
  - Not applicable. 0 day Zero days

Ihon alle:

- **Nauta**
  - Not applicable. 0 day Zero days

- **Lammas**
  - Not applicable. 0 day Zero days

- **Sika**
  - Not applicable. 0 day Zero days

Lihakseen:

- **Nauta**
  - Not applicable. 0 day Zero days

- **Lammas**
  - Not applicable. 0 day Zero days

- **Sika**
  - Not applicable. 0 day Zero days

Anatomis-terapeuttis-kemiallinen (ATCvet) luokitus:
QI02AA04

Reseptistatus:

Saatavissa vain kielillä German Estonian Greek English Italian Portuguese Norwegian

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Myyntiluvan tila:

Peruutettu

Myönnetty:

Saatavissa vain kielillä Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Lisätiedot

Luvan tyyppi:

Myyntiluvan laillinen peruste:

Saatavissa vain kielillä English Italian

Myyntiluvan haltija:

Boehringer Ingelheim Vetmedica GmbH

Myyntiluvan myöntämispäivä:

15/07/2013

Erän vapauttamisesta vastaavat valmistuspaikat:

Boehringer Ingelheim Animal Health France
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Vastaava viranomainen:
European Commission

Myyntilupanumero:
Tätä tietoa ei ole saatavana tästä valmisteesta.

Myyntiluvan tilan muutoksen päivämäärä:
15/07/2013

Ilmoitukset epäillyistä haittavaikutuksista: www.adrreports.eu/vet

Asiakirjat

Combined File of all Documents

suomi (PDF)
Julkaistu: 14/03/2026
Lataa

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

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

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