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Aftovaxpur DOE (WD) (22) O Taiwan 3/97 + Asia1 Shamir

Ekki heimilað

- Foot-and-mouth disease virus, serotype A, strain O/Taiwan/3/97, Inactivated
- Foot-and-mouth disease virus, serotype Asia 1, strain Shamir, Inactivated
- Foot-and-mouth disease virus, serotype A, strain O/Taiwan/3/97, Inactivated
- Foot-and-mouth disease virus, serotype Asia 1, strain Shamir, Inactivated

Auðkenni lyfs

Heiti lyfs:
Aftovaxpur DOE (WD) (22) O Taiwan 3/97 + Asia1 Shamir

Virkt efni:
Aðeins fáanlegt í enska
Aðeins fáanlegt í enska
Aðeins fáanlegt í enska
Aðeins fáanlegt í enska

Marktegund:
Nautgripir
Sauðkind
Svín
Nautgripir
Sauðkind

Svín

Leið stjórnsýslu:
Til notkunar í vöðva
Til notkunar undir húð
Til notkunar í vöðva
Til notkunar undir húð

Upplýsingar um lyf

Virkt efni og styrkur:
Aðeins fáanlegt í enska
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Aðeins fáanlegt í enska
Presentation_strength:≥ 6 PD50 Index:11
Aðeins fáanlegt í enska
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Aðeins fáanlegt í enska
Presentation_strength:≥ 6 PD50 Index:11

Lyfjaform:
Stungulyf, fleyti

Afurðanýtingafrestur eftir íkomuleið:
Til notkunar í vöðva:

- **Nautgripir**
  - Á ekki við. 0 dagar Zero days
- **Sauðkind**
  - Á ekki við. 0 dagar Zero days
- **Svín**

- Á ekki við. 0 dagar Zero days

Til notkunar undir húð:

- **Nautgripir**
  - Á ekki við. 0 dagar Zero days
- **Sauðkind**
  - Á ekki við. 0 dagar Zero days
- **Svín**
  - Á ekki við. 0 dagar Zero days

Til notkunar í vöðva:

- **Nautgripir**
  - Á ekki við. 0 dagar Zero days
- **Sauðkind**
  - Á ekki við. 0 dagar Zero days
- **Svín**
  - Á ekki við. 0 dagar Zero days

Til notkunar undir húð:

- **Nautgripir**
  - Á ekki við. 0 dagar Zero days
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Sauðkind

- Á ekki við. 0 dagar Zero days

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Svín

- Á ekki við. 0 dagar Zero days

ATC flokkun (dýralyf) (e. Anatomical therapeutic chemical veterinary - ATC-VET):
QI02AA04

Lögformleg staða:
Ávísunarskylt dýralyf

Staða leyfis:
Dregið til baka af leyfishafa

Heimilað í:
Austurríki , Belgía , Búlgaría , Króatía , Kýpur , Tékkland , Danmörk , Eistland , Finnland , Frakkland , Þýskaland , Grikkland , Ungverjaland , Ísland , Írland , Ítalía , Lettland , Liechtenstein , Litáen , Lúxemborg , Malta , Holland , Noregur , Pólland , Portúgal , Rúmenía , Slóvakía , Slóvenía , Spánn , Svíþjóð , Bretland (Norður-Írland)

Aðrar upplýsingar

Réttindategund:

Lagalegur grundvöllur markaðsleyfis:
Aðeins fáanlegt í enska ítalska

Markaðsleyfishafi:
Boehringer Ingelheim Vetmedica GmbH

Dagsetning markaðsleyfis:
15/07/2013

Framleiðslustaður fyrir losun lotu:
Boehringer Ingelheim Animal Health France

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Ábyrgt yfirvald:
European Commission

Markaðsleyfisnúmer:
Þessar upplýsingar eru ekki aðgengilegar fyrir þetta lyf.

Dagsetning á breytingu stöðu:
15/07/2013

Til að tilkynna upplýsingar um aukaverkanir vegna dýralyfja vinsamlegast farið á www.adrreports.eu/vet

Skjöl

Combined File of all Documents

íslenska (PDF)

Birt: 14/03/2026
Sækja

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

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