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Aftovaxpur DOE (WD) (19) O Taiwan 3/97 + A22 Iraq

Ni odobreno

- Foot-and-mouth disease virus, serotype A, strain O/Taiwan/3/97, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A22 Iraq, Inactivated
- Foot-and-mouth disease virus, serotype A, strain O/Taiwan/3/97, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A22 Iraq, Inactivated

Informacije o zdravilu

Ime zdravila:
Aftovaxpur DOE (WD) (19) O Taiwan 3/97 + A22 Iraq

Učinkovina:
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

Ciljne živalske vrste:
govedo
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
govedo
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

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Pot uporabe:
intramuskularna uporaba
Subkutana uporaba
intramuskularna uporaba
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Na voljo samo v English
Presentation_strength:≥ 6 PD50 Index:11
Na voljo samo v English
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Na voljo samo v English
Presentation_strength:≥ 6 PD50 Index:11

Farmacevtska oblika:
Emulzija za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **govedo**

- Not applicable. 0 day Zero days

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

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

Subkutana uporaba:

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

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

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

intramuskularna uporaba:

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

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

- **Pig**

- Not applicable. 0 day Zero days

Subkutana uporaba:

- **govedo**
  - Not applicable. 0 day Zero days
- **Sheep**
  - Not applicable. 0 day Zero days
- **Pig**
  - Not applicable. 0 day Zero days

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):

QI02AA04

Pravni status za dobavo/izdajo zdravila:

Zdravilo, ki se izdaja na veterinarski recept

Status dovoljenja za promet z zdravilom:

Surrendered

Ima dovoljenje za promet v:

Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Dodatne informacije

Vrsta dovoljenja:

Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Pravna podlaga za izdajo dovoljenja za promet z zdravilom:

Na voljo samo v English Italian

Imetnik dovoljenja za promet z zdravilom:

Boehringer Ingelheim Vetmedica GmbH

Datum dovoljenja za promet z zdravilom:

15/07/2013

Proizvodna mesta, odgovorna za sproščanje serij:

Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France

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Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France

Pristojni organ:
European Commission

Številka dovoljenja za promet z zdravilom:
Ta podatek za to zdravilo ni na voljo.

Datum spremembe statusa dovoljenja za promet:
15/07/2013

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Combined File of all Documents

slovenščina (PDF)
Objavljeno na dan: 14/03/2026
Prenesi

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