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Aftovaxpur DOE (WD) (04) A22 Iraq

Nem engedélyezett

- Foot-and-mouth disease virus, serotype A, strain A22 Iraq, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A22 Iraq, Inactivated

Termék azonosítása

Készítmény neve:
Aftovaxpur DOE (WD) (04) A22 Iraq

Hatóanyag:
Csak itt érhető el English
Csak itt érhető el English

Célállat faj:
szarvasmarha
juh
sertés
szarvasmarha
juh
sertés

Alkalmazás módja:
Intramuscularis alkalmazás
Subcutan alkalmazás
Subcutan alkalmazás

Intramuscularis alkalmazás

Termékjellemzők

Hatóanyag / Hatáserősség:
Csak itt érhető el English
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Csak itt érhető el English
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0

Gyógyszerforma:
Emulziós injekció

Élelmezés-egészségügyi várakozási idő alkalmazás mód alapján:
Intramuscularis alkalmazás:

- **szarvasmarha**
  - Not applicable. 0 day Zero days
- **juh**
  - Not applicable. 0 day Zero days
- **sertés**
  - Not applicable. 0 day Zero days

Subcutan alkalmazás:

- **szarvasmarha**
  - Not applicable. 0 day Zero days
- **juh**

- Not applicable. 0 day Zero days

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

Subcutan alkalmazás:

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

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

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

Intramuscularis alkalmazás:

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

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

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

Anatómiai, terápiás és kémiai állatgyógyászati (ATCvet) kód:
QI02AA04

Rendelhetőség:

Csak itt érhető el Czech Estonian English French Italian Latvian Lithuanian Portuguese Romanian Slovenian Finnish Swedish Icelandic Norwegian

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Engedélyezési státusz:

Surrendered

Engedélyezett -ben:

Csak itt érhető el Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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További információ

Jogosultsági típus:
Csak itt érhető el English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Készítmény engedélyezésének jogalapja:
Csak itt érhető el English Italian

Forgalombahozatali engedély jogosultja:
Boehringer Ingelheim Vetmedica GmbH

A forgalombahozatali engedélyezés dátuma:
15/07/2013

Gyártási tételek felszabadításáért felelős gyártóhelyek:
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Felelős hatóság:
European Commission

Engedély száma:
Ez az információ erre a készítményre vonatkozóan nem áll rendelkezésre.

Engedélyezési státusz változásának dátuma:
15/07/2013

Az állatgyógyászati készítmények mellékhatásairól a www.adrreports.eu/vet oldalon tájékozódhat

Dokumentumok

Combined File of all Documents

magyar (PDF)
Megjelent: 14/03/2026
Letöltés

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