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Aftovaxpur DOE (WD) (15) O1 BFS + A22 Iraq

Ikke autorisert

- Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A22 Iraq, Inactivated
- Foot-and-mouth disease virus, serotype O, strain O1 BFS, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A22 Iraq, Inactivated

Produkt identifikasjon

Legemidlets navn:
Aftovaxpur DOE (WD) (15) O1 BFS + A22 Iraq

Aktiv substans virkestoff:
Bare tilgjengelig i Engelsk
Bare tilgjengelig i Engelsk
Bare tilgjengelig i Engelsk
Bare tilgjengelig i Engelsk

Målarter:
storfe
sau
gris
storfe
sau

gris

Administrasjonsvei:
Intramuskulær bruk
Subkutan bruk
Intramuskulær bruk
Subkutan bruk

Produktdetaljer

Aktiv substans og styrke:
Bare tilgjengelig i Engelsk
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Bare tilgjengelig i Engelsk
Presentation_strength:≥ 6 PD50 Index:11
Bare tilgjengelig i Engelsk
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Bare tilgjengelig i Engelsk
Presentation_strength:≥ 6 PD50 Index:11

Legemiddelform:
Injeksjonsvæske, emulsjon

Tilbakeholdelsestid etter administrasjonsvei:
Intramuskulær bruk:

- **storfe**
  - Ikke aktuelt. 0 dag Zero days
- **sau**
  - Ikke aktuelt. 0 dag Zero days
- **gris**

- Ikke aktuelt. 0 dag Zero days

Subkutan bruk:

- **storfe**
  - Ikke aktuelt. 0 dag Zero days
- **sau**
  - Ikke aktuelt. 0 dag Zero days
- **gris**
  - Ikke aktuelt. 0 dag Zero days

Intramuskulær bruk:

- **storfe**
  - Ikke aktuelt. 0 dag Zero days
- **sau**
  - Ikke aktuelt. 0 dag Zero days
- **gris**
  - Ikke aktuelt. 0 dag Zero days

Subkutan bruk:

- **storfe**
  - Ikke aktuelt. 0 dag Zero days
-

sau

- Ikke aktuelt. 0 dag Zero days

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gris

- Ikke aktuelt. 0 dag Zero days

Anatomisk terapeutisk kjemisk klassifisering for veterinærpreparater (ATCvet):
QI02AA04

Juridisk status for forsyning:
Veterinært legemiddel gjenstand for veterinær forskrivning

Status for markedsføringstillatelse:
Overgitt-frafalt

Autorisert i:
AT , BE , BG , HR , Kypros , CZ , DK , EE , FI , FR , DE , EL , HU , IS , Irland , IT , LV , LI , LT , LU , MT , NL , NO , PL , PT , RO , SI , SI , ES , SE ,

Tilleggsinformasjon

Rettighetstype:

Juridisk grunnlag for markedsføringstillatelse:
Bare tilgjengelig i Engelsk Italiensk

Innehaver av markedsføringstillatelse:
Boehringer Ingelheim Vetmedica GmbH

Markedsføringsgodkjenningsdato:
15/07/2013

Tilvirker for batchfrigivelse:
Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France

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Ansvarlig myndighet:
European Commission

Godkjenningsnummer:
Denne informasjonen er ikke tilgjengelig for dette produktet.

Status for endring av markedsføringstillatelse:
15/07/2013

Rapporter om mistenkte bivirkninger: www.adrreports.eu/vet

Dokumenter

Combined File of all Documents

norsk (PDF)

Publisert på: 14/03/2026

Nedlasting

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