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Aftovaxpur DOE (WD) (03) O Taiwan 3/97

Ej godkänd

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

Produktens identitetsbeteckning

Läkemedlets namn:
Aftovaxpur DOE (WD) (03) O Taiwan 3/97

Aktiv substans:
Finns tillgänglig endast på engelska
Finns tillgänglig endast på engelska

Djurslag:
Nöt
Får
Svin
Nöt
Får
Svin

Administreringsväg:
Intramuskulär användning
Subkutan användning
Intramuskulär användning

Subkutan användning

Ytterligare information om produkten

Aktiv substans och styrka:
Finns tillgänglig endast på engelska
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Finns tillgänglig endast på engelska
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0

Läkemedelsform:
Injektionsvätska, emulsion

Karenstid per administreringsväg:
Intramuskulär användning:

- **Nöt**
  - Not applicable. 0 dygn Zero days
- **Får**
  - Not applicable. 0 dygn Zero days
- **Svin**
  - Not applicable. 0 dygn Zero days

Subkutan användning:

- **Nöt**
  - Not applicable. 0 dygn Zero days
- **Får**

- Not applicable. 0 dygn Zero days

- **Svin**
  - Not applicable. 0 dygn Zero days

Intramuskulär användning:

- **Nöt**
  - Not applicable. 0 dygn Zero days

- **Får**
  - Not applicable. 0 dygn Zero days

- **Svin**
  - Not applicable. 0 dygn Zero days

Subkutan användning:

- **Nöt**
  - Not applicable. 0 dygn Zero days

- **Får**
  - Not applicable. 0 dygn Zero days

- **Svin**
  - Not applicable. 0 dygn Zero days

Anatomisk terapeutisk kemisk veterinärkod (ATCvet-kod):
QI02AA04

Receptstatus:
Receptbelagt veterinärmedicinskt läkemedel

Godkännandestatus:
Övergiven

Godkänd i:
Österrike , Belgien , Bulgarien , Kroatien , Cypern , Tjeckien , Danmark , Estland , Finland , Frankrike , Tyskland , Grekland , Ungern , Island , Irland , Italien , Lettland , Liechtenstein , Litauen , Luxemburg , Malta , Nederländerna , Norge , Polen , Portugal , Rumänien , Slovakien , Slovenien , Spanien , Sverige , Storbritannien (Nordirland)

Ytterligare information

Typ av berättigande:

Legal grund för produktgodkännande:
Finns tillgänglig endast på engelska italienska

Innehavare av godkännande för försäljning:
Boehringer Ingelheim Vetmedica GmbH

Godkännandedatum:
15/07/2013

Tillverkningsplatser för frisläppande av tillverkningssatser:
Boehringer Ingelheim Animal Health France
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Ansvarig myndighet:
European Commission

Godkännandenummer:
Denna information finns inte tillgänglig för detta läkemedel.

Datum för ändring av godkännandestatus:
15/07/2013

Rapporter om misstänkta biverkningar: www.adrreports.eu/vet

Dokumenter

Gemensam fil för alla dokument

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Publicerad på: 14/03/2026
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