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Aftovaxpur DOE (WD) (05) A24 Cruzeiro

Niet gemachtigd

- Foot-and-mouth disease virus, serotype A, strain A24 Cruzeiro, Inactivated
- Foot-and-mouth disease virus, serotype A, strain A24 Cruzeiro, Inactivated

Productidentificatie

Naam van het geneesmiddel:
Aftovaxpur DOE (WD) (05) A24 Cruzeiro

Werkzame stof:
Alleen beschikbaar in Engels
Alleen beschikbaar in Engels

Doeldiersoort(en):
Rund
Schaap
Varken
Rund
Schaap
Varken

Toedieningsweg:
Intramusculair gebruik
Subcutaan gebruik
Intramusculair gebruik

Subcutaan gebruik

Productgegevens

Werkzame stof / Sterkte:
Alleen beschikbaar in Engels
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Alleen beschikbaar in Engels
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0

Farmaceutische vorm:
Emulsie voor injectie

Wachttijd per toedieningsweg:
Intramusculair gebruik:

- **Rund**
  - Not applicable. 0 day Zero days
- **Schaap**
  - Not applicable. 0 day Zero days
- **Varken**
  - Not applicable. 0 day Zero days

Subcutaan gebruik:

- **Rund**
  - Not applicable. 0 day Zero days
- **Schaap**

- Not applicable. 0 day Zero days

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

Intramusculair gebruik:

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

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

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

Subcutaan gebruik:

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

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

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

Anatomisch therapeutisch chemische veterinaire classificatie (ATCvet code) :

QI02AA04

Afleverstatus:

Alleen beschikbaar in Tsjechisch Estlands Engels Frans Italiaans Letlands Litouws Portugees Roemeense Toedieningsweg Fins Zweeds IJslands Norwegian

Vergunningsstatus:

Surrendered

Toegelaten in:

Oostenrijk , België , Bulgarije , Kroatië , Cyprus , Tsjechië , Denemarken , Estland , Finland , Frankrijk , Duitsland , Griekenland , Hongarije , IJsland , Ierland , Italië , Letland , Liechtenstein , Litouwen , Luxemburg , Malta , Nederland , Noorwegen , Polen , Portugal , Roemenië , Slowakije , Slovenië , Spanje , Zweden ,

Alleen beschikbaar in Estlands Engels Frans Litouws Portugees Zweeds IJslands Norwegian

Aanvullende informatie

Type vergunning voor het in de handel:

Alleen beschikbaar in Engels Frans Kroatisch Italiaans Letlands Fins Zweeds IJslands Norwegian

Wettelijke basis productvergunning:

Alleen beschikbaar in Engels Italiaans

Handelsvergunninghouder:

Boehringer Ingelheim Vetmedica GmbH

Handelsvergunningsdatum:

15/07/2013

Locaties fabrikanten vrijgifte:

Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France
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Verantwoordelijke instantie:
European Commission

Vergunningsnummer:
Deze informatie is niet beschikbaar voor dit product.

Datum toelatingswijziging:
15/07/2013

Om bijwerkingen van diergeneesmiddelen te raadplegen, zie www.adrreports.eu/vet

Documenten

Combined File of all Documents

Nederlands (PDF)
Gepubliceerd op: 14/03/2026
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