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Aftovaxpur DOE (WD) (05) A24 Cruzeiro

No autorizado

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

Identificación del medicamento

Nombre del medicamento:
Aftovaxpur DOE (WD) (05) A24 Cruzeiro

Principio activo:
Disponible únicamente en English
Disponible únicamente en English

Especies de destino:
Bovino
Ovino
Porcino
Bovino
Ovino
Porcino

Vía de administración:
Vía intramuscular
Vía subcutánea
Vía intramuscular

Vía subcutánea

Datos del medicamento

Principio activo y concentración:
Disponible únicamente en English
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0
Disponible únicamente en English
Presentation_strength:≥ 6 PD50 Reference:Hse Index:0

Forma farmacéutica:
Emulsión inyectable

Tiempo de espera por vía de administración:
Vía intramuscular:

- **Bovino**
  - Not applicable. 0 Día Zero days
- **Ovino**
  - Not applicable. 0 Día Zero days
- **Porcino**
  - Not applicable. 0 Día Zero days

Vía subcutánea:

- **Bovino**
  - Not applicable. 0 Día Zero days
- **Ovino**

- Not applicable. 0 Día Zero days

- **Porcino**
  - Not applicable. 0 Día Zero days

Vía intramuscular:

- **Bovino**
  - Not applicable. 0 Día Zero days

- **Ovino**
  - Not applicable. 0 Día Zero days

- **Porcino**
  - Not applicable. 0 Día Zero days

Vía subcutánea:

- **Bovino**
  - Not applicable. 0 Día Zero days

- **Ovino**
  - Not applicable. 0 Día Zero days

- **Porcino**
  - Not applicable. 0 Día Zero days

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código químico anatómico-terapéutico para medicamentos veterinarios (ATCvet):

QI02AA04

Condiciones de dispensación:

Disponible únicamente en Czech Estonian English French Italian Latvian Lithuanian Portuguese Romanian Slovenian Finnish Swedish Icelandic Norwegian

Estado de la autorización:

Anulado

Autorizado en:

Austria , Bélgica , Bulgaria , Croacia , Chipre , República Checa , Dinamarca , Estonia , Finlandia , Francia , Alemania , Grecia , Hungría , Islandia , Irlanda , Italia , Letonia , Liechtenstein , Lituania , Luxemburgo , Malta , Países Bajos , Noruega , Polonia , Portugal , Rumania; Rumanía , Eslovaquia , Eslovenia , España , Suecia ,

Disponible únicamente en Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Información adicional

Tipo legal de la autorización:

Disponible únicamente en English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Base legal de la autorización del medicamento:

Disponible únicamente en English Italian

Titular de la autorización de comercialización:

Boehringer Ingelheim Vetmedica GmbH

Fecha de autorización de comercialización:

15/07/2013

Fabricante responsable de la liberación del lote:

Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France
Boehringer Ingelheim Animal Health France
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Autoridad responsable:
European Commission

Número de autorización:
No se dispone de esta información para este medicamento.

Fecha de modificación del estado de la autorización:
15/07/2013

Para consultar notificaciones de presuntos efectos adversos diríjase a www.adrreports.eu/vet

Documentos

Combined File of all Documents

Español (PDF)
Publicado el: 14/03/2026
Descargar

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