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EMDOFLUXIN 50 MG/ML SOLUTION FOR INJECTION FOR CATTLE, HORSES AND PIGS

Ima dovoljenje za promet

- Flunixin meglumine

Informacije o zdravilu

Ime zdravila:
EMDOFLUXIN 50 MG/ML SOLUTION FOR INJECTION FOR CATTLE, HORSES AND PIGS

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
govedo
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Danish German Estonian Greek English Italian Latvian Lithuanian Hungarian Romanian Swedish Icelandic

Pot uporabe:
intramuskularna uporaba
Intravenska uporaba

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
83.00 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:

Raztopina za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

- **govedo**
  - Meat and offal. 31 day
  - Milk. 36 hour
- **Pig**
  - Meat and offal. 24 day

Intravenska uporaba:

- **govedo**
  - Meat and offal. 4 day
  - Milk. 24 hour
- **Equid**
  - Meat and offal. 5 day
  - Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):

QM01AG90

Pravni status za dobavo/izdajo zdravila:
Zdravilo, ki se izdaja na veterinarski recept

Status dovoljenja za promet z zdravilom:
Valid

Ima dovoljenje za promet v:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Na voljo v:
Germany

Opis ovojnine zdravila:
Na voljo samo v French
Na voljo samo v French
Na voljo samo v French

Dodatne informacije

Vrsta dovoljenja:
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Emdoka

Datum dovoljenja za promet z zdravilom:
30/06/2020

Proizvodna mesta, odgovorna za sproščanje serij:
Produlab Pharma B.V.

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:

402685.00.00

Datum spremembe statusa dovoljenja za promet:
30/06/2020

Referenčna država članica:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Številka postopka:
FR/V/0417/001

Zadevne države članice:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

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