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HEMOSILATE 125 mg/ml solution for injection

Ima dovoljenje za promet

- Etamsilate

Informacije o zdravilu

Ime zdravila:
HEMOSILATE 125 mg/ml solution for injection

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 Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
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 Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
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 Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Pot uporabe:
Intravenska uporaba
intramuskularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
125.00 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
Intravenska uporaba:

- **govedo**
  - Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

  - Milk. 0 day

- **Sheep**
  - Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

  - Milk. 0 day

- **Goat**

- Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- Milk. 0 day

- **Pig**
  - Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- **Horse**
  - Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- Milk. 0 day

intramuskularna uporaba:

- **govedo**
  - Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- Milk. 0 day

- **Sheep**
  - Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- Milk. 0 day

- **Goat**
  - Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- Milk. 0 day

- **Pig**
  - Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

- **Horse**
  - Meat and offal. no withdrawal period

Meat and offal: Adm. IV Zero days. Adm. IM 1 day

  - Milk. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QB02BX01

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

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Na voljo v:
Denmark

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

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:
Ecuphar Veterinaria S.L.U.

Datum dovoljenja za promet z zdravilom:
13/05/2020

Proizvodna mesta, odgovorna za sproščanje serij:
Zoetis Manufacturing & Research Spain S.L.

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
62253

Datum spremembe statusa dovoljenja za promet:
13/05/2020

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

Številka postopka:
ES/V/0281/001

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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Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Povzetek glavnih značilnosti zdravila

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

Navodilo za uporabo

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

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