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Vetivex 11 (Hartmann's) solution for infusion for cattle, horses, dogs and cats

Ima dovoljenje za promet

- Sodium lactate
- Sodium chloride
- Potassium chloride
- Calcium chloride dihydrate

Informacije o zdravilu

Ime zdravila:
Vetivex 11 (Hartmann's) solution for infusion for cattle, horses, dogs and cats

Učinkovina:
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
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

Pot uporabe:
Intravenska uporaba

Podatki o zdravilu

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

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

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

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

Farmacevtska oblika:
Raztopina za infundiranje

Karenca glede na pot uporabe:
Intravenska uporaba:

- **govedo**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Horse**
  - Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QB05BB01

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:
Denmark

Opis ovojnine zdravila:
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
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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:
Dechra Regulatory B.V.

Datum dovoljenja za promet z zdravilom:
23/07/2013

Proizvodna mesta, odgovorna za sproščanje serij:
Infomed Fluids S.R.L.
Industria Farmaceutica Galenica Senese S.r.l.

Pristojni organ:
Danish Medicines Agency

Številka dovoljenja za promet z zdravilom:
50931

Datum spremembe statusa dovoljenja za promet:
23/07/2013

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

Številka postopka:
IE/V/0512/001

Zadevne države članice:
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.

Combined File of all Documents

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