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SOLUZIONE ELETTROLITICA REIDRATANTE III

Ima dovoljenje za promet

- Sodium chloride
- Potassium chloride
- Magnesium chloride hexahydrate
- Calcium chloride dihydrate
- Sodium acetate
- Sodium citrate dihydrate

Informacije o zdravilu

Ime zdravila:
SOLUZIONE ELETTROLITICA REIDRATANTE III

Učinkovina:
Na voljo samo v English
Na voljo samo v English
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
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Pot uporabe:
Intravenska uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
5.00 gram(s) / 1000.00 millilitre(s)

Na voljo samo v English
0.75 gram(s) / 1000.00 millilitre(s)

Na voljo samo v English
0.31 gram(s) / 1000.00 millilitre(s)

Na voljo samo v English
0.35 gram(s) / 1000.00 millilitre(s)

Na voljo samo v English
6.40 gram(s) / 1000.00 millilitre(s)

Na voljo samo v English
0.75 gram(s) / 1000.00 millilitre(s)

Farmacevtska oblika:
Raztopina za infundiranje

Karenca glede na pot uporabe:

Intravenska uporaba:

- **govedo**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Sheep**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **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 brez veterinarskega recepta

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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Opis ovojnine zdravila:

Na voljo samo v Italian

Na voljo samo v Italian

Na voljo samo v Italian

Na voljo samo v Italian
Na voljo samo v Italian
Na voljo samo v Italian

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:
Industria Farmaceutica Galenica Senese S.r.l.

Datum dovoljenja za promet z zdravilom:
29/09/1999

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

Pristojni organ:
Ministry Of Health

Številka dovoljenja za promet z zdravilom:
Ta podatek za to zdravilo ni na voljo.

Datum spremembe statusa dovoljenja za promet:
29/09/1999

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.