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Calcibel Forte, 380/60/50 mg/ml solution for infusion for horses, cattle, sheep, goats and pigs

Ima dovoljenje za promet

- Calcium gluconate
- Magnesium chloride hexahydrate
- Boric acid

Informacije o zdravilu

Ime zdravila:
Calcibel Forte, 380/60/50 mg/ml solution for infusion for horses, cattle, sheep, goats and pigs

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

Ciljne živalske vrste:
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
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
380.00 milligram(s) / 1.00 millilitre(s)

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

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

Farmacevtska oblika:
Raztopina za infundiranje

Karenca glede na pot uporabe:
Intravenska uporaba:

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

Sheep

- Milk. 0 hour
- Meat and offal. 0 day

•

Goat

- Milk. 0 hour
- Meat and offal. 0 day

•

Pig

- Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QA12AX

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:
Portugal

Opis ovojnine zdravila:
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:
Bela-Pharm GmbH & Co. KG

Datum dovoljenja za promet z zdravilom:
9/08/2021

Proizvodna mesta, odgovorna za sproščanje serij:
Bela-Pharm GmbH & Co. KG

Pristojni organ:
Directorate General For Food And Veterinary

Številka dovoljenja za promet z zdravilom:
1444/01/21DFVPT

Datum spremembe statusa dovoljenja za promet:
2/06/2025

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

Številka postopka:
CZ/V/0170/001

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 Greek 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

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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.