File downloaded on 2026-11-21
Source URL: https://medicines.health.europa.eu/veterinary/lv/600000053625

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

Atļautas

- Calcium gluconate
- Magnesium chloride hexahydrate
- Boric acid

Zāļu identifikācija

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

Aktīvā viela:
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English

Mērķsugas:
Zirgs
Liellops
Aita
Kaza
Cūka

Lietošanas veids:
intravenozai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

Pieejamas tikai English
380.00 miligrams(i) / 1.00 mililitrs(i)

Pieejamas tikai English
60.00 miligrams(i) / 1.00 mililitrs(i)

Pieejamas tikai English
50.00 miligrams(i) / 1.00 mililitrs(i)

Farmaceitiskā forma:

šķīdums infūzijām

Ierobežojumu periods pēc lietošanas veida:

intravenozai lietošanai:

- **Zirgs**
  - Piens. 0 stunda
  - Gaļa un blakusprodukti. 0 diena
- **Liellops**
  - Piens. 0 stunda
  - Gaļa un blakusprodukti. 0 diena
- **Aita**
  - Piens. 0 stunda
  - Gaļa un blakusprodukti. 0 diena
- **Kaza**
  - Piens. 0 stunda
  - Gaļa un blakusprodukti. 0 diena

-

Cūka

- Gaļa un blakusprodukti. 0 diena

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QA12AX

Izplatīšanas juridiskie nosacījumi:
Recepšu veterinārās zāles

Atļaujas statuss:
Derīga

Atļautas:
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Pieejams:
Greece

Iepakojuma apraksts:
Pieejamas tikai English
Pieejamas tikai English

Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:
Pieteikums ģenēriskām veterinārajām zālēm (Direktīvas 2001/82/EC 13.(1) pants)

Tirdzniecības atļaujas turētājs:
Bela-Pharm GmbH & Co. KG

Tirdzniecības atļaujas datums:
20/09/2021

Ražošanas vietas sēriju izlaidei:
Bela-Pharm GmbH & Co. KG

Atbildīgā iestāde:
National Organization For Medicines

Atļaujas numurs:
84808/21-09-2021/K-0214302

Atļaujas statusa maiņas datums:
20/09/2021

Atsauces dalībvalsts:
Čehija

Procedūras numurs:
CZ/V/0170/001

Attiecīgās dalībvalstis:
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian Greek English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish German Estonian English French Italian Dutch Portuguese Slovak Finnish Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Italian Lithuanian Dutch Portuguese Romanian Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

---

Lai uzzinātu par veterināro zāļu blakusparādībām, lūdzu, dodieties uz: www.adrreports.eu/vet

Dokumenti

Zāļu apraksts

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.

Lietošanas instrukcija

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.

Marķējuma teksts

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.