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Source URL: https://medicines.health.europa.eu/veterinary/sl/600000092620
Kaltetan, 250 mg/ml + 80 mg/ml + 10 mg/ml solution for infusion for horses, cattle and pigs

- CALCIUM GLUCONATE FOR INJECTION
- Magnesium chloride hexahydrate
- Sodium glycerophosphate pentahydrate

Ima dovoljenje za promet

Informacije o zdravilu

Ime zdravila:
Kaltetan, 250 mg/ml + 80 mg/ml + 10 mg/ml solution for infusion for horses, cattle and pigs
U?inkovina:

- 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
- Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
- govedo

Pot uporabe:

- Intravenska uporaba

Podatki o zdravilu

U?inkovina / Jakost:

- Na voljo samo v English
  250.00
  milligram(s)/millilitre
  /
  1.00
  milligram(s)/millilitre
- Na voljo samo v English
  80.00
  milligram(s)
  /
  1.00
  millilitre(s)
- Na voljo samo v English

10.00
milligram(s)
/
1.00
millilitre(s)

Farmacevtska oblika:

- Raztopina za infundiranje

Karenca glede na pot uporabe:

- Intravenska uporaba
  - Horse
    - Milk
      5
      week
      Meat and offal: Zero days. Milk: Zero hours
  - Pig
    - Meat and offal
      5
      week
      Meat and offal: Zero days
  - govedo
    - Milk
      5
      week
      Meat and offal: Zero days. Milk: Zero hours

Anatomsko-terapevtsko-kemi?na 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 Lithuanian Dutch Portuguese Romanian Slovak Swedish Icelandic Norwegian

Na voljo v:

- Romania

Opis ovojnine zdravila:

- 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:

- Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Datum dovoljenja za promet z zdravilom:

- 15/03/2022

Proizvodna mesta, odgovorna za sproš?anje serij:

- Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Pristojni organ:

- Institute For Control Of Biological Products And Veterinary Medicines

Številka dovoljenja za promet z zdravilom:

- 220055

Datum spremembe statusa dovoljenja za promet:

- 10/06/2025

Referen?na država ?lanica:

- Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Številka postopka:

- PL/V/0110/001

Zadevne države ?lanice:

- 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 Latvian Dutch Portuguese Slovak Swedish Icelandic Norwegian
- 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 Spanish Czech German Estonian English French Italian Lithuanian Dutch Portuguese Romanian Slovak Swedish Icelandic Norwegian
- Na voljo samo v Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Za poro?ila o domnevnih neželenih u?inkih zdravil za uporabo v veterinaski medicini obiš?ite stran: www.adrreports.eu/vet

Dokumenti

Informacije o zdravilu

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenš?ina). Lahko ga najdete spodaj v drugem jeziku.
Drugi jeziki (1)
Romanian (PDF)
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