NARKAMON 50 mg/ml injekčný roztok

Ima dovoljenje za promet

- Ketamine hydrochloride

Informacije o zdravilu

Ime zdravila:
NARKAMON 50 mg/ml injekčný roztok

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
tele
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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Icelandic Norwegian
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Pot uporabe:
intramuskularna uporaba
Intraperitonealna uporaba
Intravenska uporaba

Podatki o zdravilu

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

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **tele**
  - Meat and offal. 24 hour
- **Sheep**
  - Meat and offal. 24 hour
  - Milk. 0 day milk zero days
- **Goat**
  - Meat and offal. 24 hour
  - Milk. 0 day milk zero days
- **Dog**
  - Not applicable. 0 day Not applicable
- **Cat**
  - Not applicable. 0 day Not applicable
- **Guinea pig**
  - Not applicable. 0 day Not applicable
- **Monkey**
  - Not applicable. 0 day Not applicable

- **Parrot**
  - Not applicable. 0 day Not applicable
- **plazilec**
  - Not applicable. 0 day Not applicable
- **Antelope**
  - Meat and offal. 24 hour
- **Roe deer**
  - Meat and offal. 24 hour
- **Pigeon**
  - Meat and offal. 24 hour
- **Red deer**
  - Meat and offal. 24 hour

Intraperitonealna uporaba:

- **Mouse**
  - Not applicable. 0 day Not applicable
- **Rat**
  - Not applicable. 0 day Not applicable

Intravenska uporaba:

- **Horse**

- Meat and offal. 24 hour
- Milk. 0 day milkmzreo days

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Dog

- Not applicable. 0 day Not applicable

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QN01AX03

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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Opis ovojnine zdravila:
Na voljo samo v Slovak

Dodatne informacije

Vrsta dovoljenja:
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English French Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Bioveta a.s.

Datum dovoljenja za promet z zdravilom:

26/09/1988

Proizvodna mesta, odgovorna za sproščanje serij:
Bioveta a.s.

Pristojni organ:
Institute For State Control Of Veterinary Biologicals And Medicaments

Številka dovoljenja za promet z zdravilom:
99/172/88-C/S

Datum spremembe statusa dovoljenja za promet:
26/09/1988

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.