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Source URL: https://medicines.health.europa.eu/veterinary/sl/600000093171

Ketexx 100 mg/ml solution for injection

Ima dovoljenje za promet

- Ketamine hydrochloride

Informacije o zdravilu

Ime zdravila:
Ketexx 100 mg/ml solution for injection

Učinkovina:
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 Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
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Pot uporabe:
intramuskularna uporaba
Intravenska uporaba
Intraperitonealna uporaba

Podatki o zdravilu

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

Farmacevtska oblika:
Raztopina za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **govedo**
  - Meat and offal. 1 day
  - Milk. 0 day

- **Rabbit (exclusively kept as pet)**
  - All relevant tissues. no withdrawal period

Not authorised for use in rabbits for human consumption.

- **Horse**
  - Meat and offal. 1 day
  - Milk. 0 day

Intravenska uporaba:

- **govedo**
  - Meat and offal. 1 day
  - Milk. 0 day

- **Rabbit (exclusively kept as pet)**
  - All relevant tissues. no withdrawal period

Not authorised for use in rabbits for human consumption.

- **Horse**
  - Meat and offal. 1 day
  - Milk. 0 day

- **Sheep**
  - Meat and offal. 1 day
  - Milk. 0 day

- **Goat**
  - Meat and offal. 1 day
  - Milk. 0 day

Intraperitonealna uporaba:

- **Rabbit (exclusively kept as pet)**
  - All relevant tissues. no withdrawal period

Not authorised for use in rabbits for human consumption.

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 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
Na voljo samo v English
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

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

Imetnik dovoljenja za promet z zdravilom:
Alfasan Nederland B.V.

Datum dovoljenja za promet z zdravilom:
20/05/2022

Proizvodna mesta, odgovorna za sproščanje serij:
Alfasan Nederland B.V.

Pristojni organ:
Health Products Regulatory Authority

Številka dovoljenja za promet z zdravilom:
VPA10980/016/001

Datum spremembe statusa dovoljenja za promet:
20/05/2022

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

Številka postopka:
FR/V/0435/001

Zadevne države članice:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak 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.

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