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Ketexx 100 mg/ml solution for injection

Atļautas

- Ketamine hydrochloride

Zāļu identifikācija

Zāļu nosaukums:
Ketexx 100 mg/ml solution for injection

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Liellops
Žurka
Pele
Kāmis
Jūrascūciņa
Trusis (lolojumdzīvnieks)
Kaķis
Zirgs
Suns
Aita
Kaza

Lietošanas veids:
intramuskulārai lietošanai
intravenozai lietošanai

intraperitoneālai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

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

Farmaceitiskā forma:

Šķīdums injekcijām

Ierobežojumu periods pēc lietošanas veida:

intramuskulārai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 1 diena
  - Piens. 0 diena
- **Trusis (lolojumdzīvnieks)**
  - Visi attiecīgie audi. no withdrawal period

Not authorised for use in rabbits for human consumption.

- **Zirgs**
  - Gaļa un blakusprodukti. 1 diena
  - Piens. 0 diena

intravenozai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 1 diena
  - Piens. 0 diena
- **Trusis (lolojumdzīvnieks)**

- Visi attiecīgie audi. no withdrawal period

Not authorised for use in rabbits for human consumption.

- **Zirgs**
  - Gaļa un blakusprodukti. 1 diena
  - Piens. 0 diena
- **Aita**
  - Gaļa un blakusprodukti. 1 diena
  - Piens. 0 diena
- **Kaza**
  - Gaļa un blakusprodukti. 1 diena
  - Piens. 0 diena

intraperitoneālai lietošanai:

- **Trusis (lolojumdzīvnieks)**
  - Visi attiecīgie audi. no withdrawal period

Not authorised for use in rabbits for human consumption.

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QN01AX03

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

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Iepakojuma apraksts:
Pieejamas tikai English
Pieejamas tikai English
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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:
Alfasan Nederland B.V.

Tirdzniecības atļaujas datums:
20/05/2022

Ražošanas vietas sēriju izlaidei:
Alfasan Nederland B.V.

Atbildīgā iestāde:
Health Products Regulatory Authority

Atļaujas numurs:
VPA10980/016/001

Atļaujas statusa maiņas datums:
20/05/2022

Atsauces dalībvalsts:

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

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Procedūras numurs:
FR/V/0435/001

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

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

Dokumenti

Visu dokumentu apvienotais fails

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

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