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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:
29/06/2022

Ražošanas vietas sēriju izlaidei:
Alfasan Nederland B.V.

Atbildīgā iestāde:
Danish Medicines Agency

Atļaujas numurs:
65647

Atļaujas statusa maiņas datums:
29/06/2022

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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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.

Zāļu apraksts

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

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