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KETABEL 100 MG/ML SOLUTION FOR INJECTION

Atļautas

- Ketamine hydrochloride

Zāļu identifikācija

Zāļu nosaukums:
KETABEL 100 MG/ML SOLUTION FOR INJECTION

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Liellops
Cūka
Žurka
Jūrascūciņa
Trusis
Kaķis
Aita
Kaza
Suns
Zirgs
Kāmis
Pele

Lietošanas veids:
intramuskulārai lietošanai

intravenozai lietošanai
intraperitoneālai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:
Pieejamas tikai English
115.34 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
- **Cūka**
  - Gaļa un blakusprodukti. 1 diena
- **Aita**
  - Gaļa un blakusprodukti. 1 diena
  - Piens. 0 diena
- **Kaza**
  - Gaļa un blakusprodukti. 1 diena
  - Piens. 0 diena

intravenozai lietošanai:

- **Liellops**

- Gaļa un blakusprodukti. 1 diena

- Piens. 0 diena

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Zirgs

- Gaļa un blakusprodukti. 1 diena

- Piens. 0 diena

•

Aita

- Gaļa un blakusprodukti. 1 diena

- Piens. 0 diena

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Kaza

- Gaļa un blakusprodukti. 1 diena

- Piens. 0 diena

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 French
Pieejamas tikai French
Pieejamas tikai French
Pieejamas tikai French

Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:
Pieteikums ģenēriskām veterinārajām zālēm (Direktīvas 2001/82/EC 13.(1) pants)

Tirdzniecības atļaujas turētājs:
Bela-Pharm GmbH & Co. KG

Tirdzniecības atļaujas datums:
11/06/2020

Ražošanas vietas sēriju izlaidei:
Bela-Pharm GmbH & Co. KG

Atbildīgā iestāde:
French Agency For Food, Environmental And Occupational Health & Safety

Atļaujas numurs:
FR/V/2514272 5/2020

Atļaujas statusa maiņas datums:
10/03/2023

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

Procedūras numurs:
FR/V/0338/001

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

Zāļu apraksts

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

Visu dokumentu apvienotais fails

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

Apvienotais marķējuma un lietošanas instrukcijas teksts

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

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