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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

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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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Pieejams:
Denmark

Iepakojuma apraksts:
Pieejamas tikai English
Pieejamas tikai English
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:
Bela-Pharm GmbH & Co. KG

Tirdzniecības atļaujas datums:
25/04/2022

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

Atbildīgā iestāde:
Danish Medicines Agency

Atļaujas numurs:
66501

Atļaujas statusa maiņas datums:
25/04/2022

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

Visu dokumentu apvienotais fails

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

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