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MENBUTIL 100 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS, HORSES, SHEEP AND GOATS

Atļautas

- Menbutone

Zāļu identifikācija

Zāļu nosaukums:
MENBUTIL 100 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS, HORSES, SHEEP AND GOATS

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Liellops (teļš)
Aita
Kaza
Liellops
Cūka
Zirgs

Lietošanas veids:
intramuskulārai lietošanai
intravenozai lietošanai
ievadīšanai tesmenī

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

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

Farmaceitiskā forma:

Šķīdums injekcijām

Ierobežojumu periods pēc lietošanas veida:

intramuskulārai lietošanai:

- **Liellops (teļš)**
  - Gaļa un blakusprodukti. 0 diena
- **Aita**
  - Gaļa un blakusprodukti. 0 diena
  - Piens. 0 diena
- **Kaza**
  - Gaļa un blakusprodukti. 0 diena
  - Piens. 0 diena

intravenozai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 0 diena
  - Piens. 0 diena
- **Cūka**
  - Gaļa un blakusprodukti. 0 diena
-

Zirgs

- Gaļa un blakusprodukti. 0 diena

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Aita

- Gaļa un blakusprodukti. 0 diena
- Piens. 0 diena

•

Kaza

- Gaļa un blakusprodukti. 0 diena
- Piens. 0 diena

ievadīšanai tesmenī:

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Cūka

- Gaļa un blakusprodukti. 0 diena

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QA05AX90

Izplatīšanas juridiskie nosacījumi:
Recepšu veterinārās zāles

Atļaujas statuss:
Derīga

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

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Pieejams:
Cyprus

Iepakojuma apraksts:
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:
aniMedica GmbH

Tirdzniecības atļaujas datums:
18/11/2018

Ražošanas vietas sēriju izlaidei:
aniMedica GmbH
Industrial Veterinaria S.A.

Atbildīgā iestāde:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Atļaujas numurs:
CY00709V

Atļaujas statusa maiņas datums:
11/01/2022

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

Procedūras numurs:
FR/V/0200/001

Attiecīgās dalībvalstis:
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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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