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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 English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Pieejams:
Sweden

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:
15/01/2019

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

Atbildīgā iestāde:
Swedish Medical Products Agency

Atļaujas numurs:
58074

Atļaujas statusa maiņas datums:
15/01/2019

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:
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

Lietošanas instrukcija

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.

Marķējuma teksts

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.

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