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Ketink 100 mg/ml solution for injection for cattle, pigs and horses

Atļautas

- Ketoprofen

Zāļu identifikācija

Zāļu nosaukums:
Ketink 100 mg/ml solution for injection for cattle, pigs and horses

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Liellops
Cūka
Zirgs

Lietošanas veids:
intramuskulārai lietošanai
intravenozai lietošanai

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**
  - Gaļa un blakusprodukti. 4 diena
  - Piens. 0 stunda
- **Cūka**
  - Gaļa un blakusprodukti. 4 diena
- **Liellops**
  - Gaļa un blakusprodukti. 4 diena
  - Piens. 0 stunda

intravenozai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 4 diena
  - Piens. 0 stunda
- **Zirgs**
  - Gaļa un blakusprodukti. 4 diena
  - Piens. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

- **Liellops**
  - Gaļa un blakusprodukti. 4 diena
  - Piens. 0 stunda

-

Zirgs

- Gaļa un blakusprodukti. 4 diena
- Piens. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QM01AE03

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
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:
Industrial Veterinaria S.A.

Tirdzniecības atļaujas datums:
27/02/2012

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

Atbildīgā iestāde:
Danish Medicines Agency

Atļaujas numurs:
48707

Atļaujas statusa maiņas datums:
27/02/2012

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

Procedūras numurs:
ES/V/0175/001

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

Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish German Estonian English French Italian Dutch Portuguese Slovak Finnish Swedish Icelandic Norwegian
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Pieejamas tikai Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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.

Lietošanas instrukcija

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.