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Loxicom 20 mg/ml - Solution for injection (cattle, pigs, horses)

Atļautas

- Meloxicam

Zāļu identifikācija

Zāļu nosaukums:
Loxicom 20 mg/ml - Solution for injection (cattle, pigs, horses)

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Liellops
Zirgs
Cūka

Lietošanas veids:
intramuskulārai lietošanai
intravenozai lietošanai
subkutānai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:
Pieejamas tikai English
Presentation_strength:20 mg Reference:B P Index:0

Farmaceitiskā forma:
Šķīdums injekcijām

Ierobežojumu periods pēc lietošanas veida:
intramuskulārai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 15 diena 15 days
  - Milk. 5 diena 5 days
- **Zirgs**
  - Gaļa un blakusprodukti. 5 diena 5 days
- **Cūka**
  - Gaļa un blakusprodukti. 5 diena 5 days

intravenozai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 15 diena 15 days
  - Milk. 5 diena 5 days
- **Zirgs**
  - Gaļa un blakusprodukti. 5 diena 5 days
- **Cūka**
  - Gaļa un blakusprodukti. 5 diena 5 days

subkutānai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 15 diena 15 days
  - Milk. 5 diena 5 days
- **Zirgs**
  - Gaļa un blakusprodukti. 5 diena 5 days
- **Cūka**
  - Gaļa un blakusprodukti. 5 diena 5 days

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QM01AC06

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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Pieejamas tikai Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Pieejams:
Belgium , Cyprus , Denmark , Finland , Germany , Greece , Ireland , Italy , Luxembourg , Netherlands , Poland , Portugal , Romania , Spain , Sweden ,
United Kingdom (Northern Ireland) , Čehija

Iepakojuma apraksts:

Pieejamas tikai English

Pieejamas tikai English

Pieejamas tikai English

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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:
Norbrook Laboratories (Ireland) Limited

Tirdzniecības atļaujas datums:
10/02/2009

Ražošanas vietas sēriju izlaidei:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Atbildīgā iestāde:
European Commission

Atļaujas numurs:
Šī informācija par šīm zālēm nav pieejama.

Atļaujas statusa maiņas datums:
13/07/2011

Lai uzzinātu par veterināro zāļu blakusparādībām, lūdzu, dodieties uz:
www.adrreports.eu/vet

Dokumenti

Visu dokumentu apvienotais fails

latviešu (PDF)
Publicēts: 14/03/2026
Lejupielādēt

ema-puar-loxicom-v-141-var-x-0003-en.pdf

ema-puar-loxicom-v-141-var-x-0005-en.pdf

ema-puar-loxicom-v-141-par-en.pdf

ema-puar-loxicom-v-141-var-x-0009-en.pdf