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Melovem 5 mg/ml - Solution for injection (Cattle, Pig)

Atļautas

- Meloxicam

Zāļu identifikācija

Zāļu nosaukums:
Melovem 5 mg/ml - Solution for injection (Cattle, Pig)

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Liellops
Cūka

Lietošanas veids:
intramuskulārai lietošanai
subkutānai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:
Pieejamas tikai English
5.00 miligrams(i) / 1.00 mililitrs(i)

Farmaceitiskā forma:

Šķīdums injekcijām

Ierobežojumu periods pēc lietošanas veida:
intramuskulārai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 15 diena

15 days. Not authorised for use in animals producing milk for human consumption.

- **Cūka**
  - Gaļa un blakusprodukti. 5 diena 5 days

subkutānai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 15 diena

15 days. Not authorised for use in animals producing milk for human consumption.

- **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

Pieejams:
Austria , Belgium , Bulgaria , Cyprus , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland) , Čehija

Iepakojuma apraksts:
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:
Dopharma Research B.V.

Tirdzniecības atļaujas datums:
7/07/2009

Ražošanas vietas sēriju izlaidei:
Dopharma B.V.

Atbildīgā iestāde:
European Commission

Atļaujas numurs:
Šī informācija par šīm zālēm nav pieejama.

Atļaujas statusa maiņas datums:
7/07/2009

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-v0152-melovem-vra-0015-en.pdf

ema-puar-melovem-v-152-par-en.pdf

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