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Source URL: https://medicines.health.europa.eu/veterinary/en/600000017668

Labiprofen 150 mg/ml solution for injection

Atļautas

- Ketoprofen

Zāļu identifikācija

Zāļu nosaukums:
Labiprofen 150 mg/ml solution for injection

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Cūka
Liellops
Zirgs

Lietošanas veids:
intramuskulārai lietošanai
intravenozai lietošanai

Sīkāka informācija par zālēm

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

Farmaceitiskā forma:
Šķīdums injekcijām

Ierobežojumu periods pēc lietošanas veida:
intramuskulārai lietošanai:

- **Cūka**
  - Gaļa un blakusprodukti. 3 diena
- **Liellops**
  - Gaļa un blakusprodukti. 2 diena
  - Piens. 0 stunda

intravenozai lietošanai:

- **Zirgs**
  - Gaļa un blakusprodukti. 1 diena
  - Piens. no withdrawal period

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

- **Liellops**
  - Gaļa un blakusprodukti. 2 diena
  - Piens. 0 stunda

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

Pieejams:
France

Iepakojuma apraksts:
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English
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:
Labiana Life Sciences S.A.

Tirdzniecības atļaujas datums:
24/06/2021

Ražošanas vietas sēriju izlaidei:
Labiana Life Sciences S.A.

Atbildīgā iestāde:
French Agency For Food, Environmental And Occupational Health & Safety

Atļaujas numurs:
FR/V/5860082 9/2021

Atļaujas statusa maiņas datums:
24/06/2021

Atsauces dalībvalsts:

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

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Procedūras numurs:
ES/V/0388/001

Attiecīgās dalībvalstis:
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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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.

Apvienotais marķējuma un lietošanas instrukcijas teksts

Šis dokuments neeksistē šādā valodā(latviešu). Jūs to varat atrast citā valodā zemāk.