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Ecomectin 10 mg/ml Solution for Injection

Atļautas

- Ivermectin

Zāļu identifikācija

Zāļu nosaukums:
Ecomectin 10 mg/ml Solution for Injection

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Liellops
Aita
Cūka

Lietošanas veids:
subkutānai lietošanai

Sīkāka informācija par zālēm

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

Farmaceitiskā forma:

Šķīdums injekcijām

Ierobežojumu periods pēc lietošanas veida:
subkutānai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 49 diena
  - Piens. no withdrawal period

Do not use in lactating cows producing milk for human consumption. Do not use in non-lactating dairy cows, including pregnant dairy heifers, within 60 days of calving.

- **Aita**
  - Gaļa un blakusprodukti. 42 diena
  - Piens. no withdrawal period

Do not use in lactating ewes producing milk for human consumption. Do not use in sheep which are intended to produce milk for human consumption within 60 days of lambing.

- **Cūka**
  - Gaļa un blakusprodukti. 28 diena

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QP54AA01

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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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 vispāratzītām veterinārajām zālēm (Direktīvas 2001/82/EC 13.(a) pants)

Tirdzniecības atļaujas turētājs:
Eco Animal Health Europe Limited

Tirdzniecības atļaujas datums:
5/02/2004

Ražošanas vietas sēriju izlaidei:
Produlab Pharma B.V.
Divasa Farmavic S.A.

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

Atļaujas numurs:
FR/V/4934184 5/2004

Atļaujas statusa maiņas datums:
5/02/2009

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

Procedūras numurs:
IE/V/0144/001

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

Apvienotais marķējuma un lietošanas instrukcijas teksts

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