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MACROSYN 100 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS AND SHEEP

Atļautas

- Tulathromycin

Zāļu identifikācija

Zāļu nosaukums:
MACROSYN 100 MG/ML SOLUTION FOR INJECTION FOR CATTLE, PIGS AND SHEEP

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Aita
Cūka
Liellops

Lietošanas veids:
intramuskulārai lietošanai
subkutānai 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:

- **Aita**
  - Gaļa un blakusprodukti. 16 diena
  - Piens. no withdrawal period

Not authorised for use in animals producing milk for human consumption. Do not use in pregnant animals, which are intended to produce milk for human consumption, within 2 months of expected parturition.

- **Cūka**
  - Gaļa un blakusprodukti. 13 diena

subkutānai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 22 diena
  - Piens. no withdrawal period

Not authorised for use in animals producing milk for human consumption. Do not use in pregnant animals, which are intended to produce milk for human consumption, within 2 months of expected parturition.

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QJ01FA94

Izplatīšanas juridiskie nosacījumi:
Recepšu veterinārās zāles

Atļaujas statuss:
Derīga

Atļautas:
Pieejamas tikai Spanish German Estonian English French Italian Dutch Portuguese Slovak Finnish Swedish Icelandic Norwegian

Iepakojuma apraksts:
Pieejamas tikai French
Pieejamas tikai French
Pieejamas tikai French
Pieejamas tikai French

Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:
Pieteikums ģenēriskām veterinārajām zālēm (Direktīvas 2001/82/EC 13.(1) pants)

Tirdzniecības atļaujas turētājs:
Bimeda Animal Health Limited

Tirdzniecības atļaujas datums:
21/09/2020

Ražošanas vietas sēriju izlaidei:
Bimeda Animal Health Limited

Atbildīgā iestāde:
State Agency Of Medicines

Atļaujas numurs:
2254

Atļaujas statusa maiņas datums:
21/09/2020

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

Procedūras numurs:
FR/V/0418/001

Attiecīgās dalībvalstis:
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak 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.

Visu dokumentu apvienotais fails

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

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