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Tulissin 100 mg/ml - Solution for injection (cattle, pigs, sheep)

Atļautas

- Tulathromycin

Zāļu identifikācija

Zāļu nosaukums:
Tulissin 100 mg/ml - Solution for injection (cattle, pigs, sheep)

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Liellops
Aita
Cūka

Lietošanas veids:
intramuskulārai lietošanai
subkutānai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:
Pieejamas tikai English
100.00 miligrams(i) / 1.00 Flakons

Farmaceitiskā forma:
Šķīdums injekcijām

Ierobežojumu periods pēc lietošanas veida:
intramuskulārai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 22 diena 22 days
- **Aita**
  - Gaļa un blakusprodukti. 16 diena 16 days
- **Cūka**
  - Gaļa un blakusprodukti. 13 diena 13 days

subkutānai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 22 diena 22 days
- **Aita**
  - Gaļa un blakusprodukti. 16 diena 16 days
- **Cūka**
  - Gaļa un blakusprodukti. 13 diena 13 days

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QJ01FA94

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 , Croatia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Netherlands , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , United Kingdom (Northern Ireland) , Čehija

Iepakojuma apraksts:
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:
Virbac

Tirdzniecības atļaujas datums:
24/04/2020

Ražošanas vietas sēriju izlaidei:
Fareva Amboise
Virbac

Atbildīgā iestāde:
European Commission

Atļaujas numurs:
Šī informācija par šīm zālēm nav pieejama.

Atļaujas statusa maiņas datums:
24/04/2020

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-tulissin-v-5073-par-en.pdf