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Zeleris 400 mg/ml florfenicol + 5 mg/ml meloxicam - Solution for injection

Atļautas

- Florfenicol
- Meloxicam

Zāļu identifikācija

Zāļu nosaukums:
Zeleris 400 mg/ml florfenicol + 5 mg/ml meloxicam - Solution for injection

Aktīvā viela:
Pieejamas tikai English
Pieejamas tikai English

Mērķsugas:
Liellops

Lietošanas veids:
subkutānai lietošanai

Sīkāka informācija par zālēm

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

Pieejamas tikai English
5.00 miligrams(i) / 1.00 mililitrs(i)

Farmaceitiskā forma:

Šķīdums injekcijām

Ierobežojumu periods pēc lietošanas veida:

subkutānai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 56 diena 56 days
  - Piens. no withdrawal period

999 days - Not for use in animals from which milk is produced for human consumption.

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:

QJ01BA99

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

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

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Iepakojuma apraksts:

Pieejamas tikai English

Pieejamas tikai English

Pieejamas tikai English

Papildu informācija

Tiesību tips:

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

Tirdzniecības atļaujas turētājs:
Ceva Sante Animale

Tirdzniecības atļaujas datums:
15/05/2017

Ražošanas vietas sēriju izlaidei:
Ceva Sante Animale

Atbildīgā iestāde:
European Commission

Atļaujas numurs:
Šī informācija par šīm zālēm nav pieejama.

Atļaujas statusa maiņas datums:
15/05/2017

Lai uzzinātu par veterināro zāļu blakusparādībām, lūdzu, dodieties uz: www.adrreports.eu/vet

Dokumenti

Visu dokumentu apvienotais fails

latviešu (PDF)

Publicēts: 13/03/2026

Lejupielādēt

ema-puar-v4099-zeleris-vra-0005-g-en.pdf

ema-puar-zeleris-v-4099-par-en.pdf