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Spirovet 600 000 iu/ml solution for injection for cattle

Atļautas

- Spiramycin

Zāļu identifikācija

Zāļu nosaukums:
Spirovet 600 000 iu/ml solution for injection for cattle

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Liellops

Lietošanas veids:
intramuskulārai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:
Pieejamas tikai English
600000.00 starptautiskā(s) vienība(s) / 1.00 mililitrs(i)

Farmaceitiskā forma:
Šķīdums injekcijām

Ierobežojumu periods pēc lietošanas veida:

intramuskulārai lietošanai:

- **Liellops**

  - Gaļa un blakusprodukti. 75 diena

Mastitis (30 000 IU of spiramycin per kg bodyweight (i.e. 5 ml of product per 100 kg bodyweight) twice at 24h of interval).

  - Piens. no withdrawal period

Respiratory infections (100 000 IU of spiramycin per kg bodyweight (i.e. 5 ml of product per 30 kg bodyweight) twice at 48h of interval). In case of treatment at the dose required for respiratory diseases, the veterinary medicinal product is not authorised for use in animals producing milk for human consumption.

  - Gaļa un blakusprodukti. 75 diena

Respiratory infections (100 000 IU of spiramycin per kg bodyweight (i.e. 5 ml of product per 30 kg bodyweight) twice at 48h of interval).

  - Piens. 14 diena

Mastitis (30 000 IU of spiramycin per kg bodyweight (i.e. 5 ml of product per 100 kg bodyweight) twice at 24h of interval).

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QJ01FA02

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 ģenēriskām veterinārajām zālēm (Direktīvas 2001/82/EC 13.(1) pants)

Tirdzniecības atļaujas turētājs:
Ceva Sante Animale

Tirdzniecības atļaujas datums:
9/11/2012

Ražošanas vietas sēriju izlaidei:
Ceva Sante Animale

Atbildīgā iestāde:
Health Products Regulatory Authority

Atļaujas numurs:
VPA10815/021/001

Atļaujas statusa maiņas datums:
9/11/2012

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

Procedūras numurs:
FR/V/0244/001

Attiecīgās dalībvalstis:

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

Visu dokumentu apvienotais fails

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

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