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Evalon (--) - Suspension and solvent for oral spray

Atļautas

- Eimeria acervulina, strain 003, Live
- Eimeria maxima, strain 013, Live
- Eimeria necatrix, strain 033, Live
- Eimeria brunetti, strain 034, Live
- Eimeria tenella, strain 004, Live

Zāļu identifikācija

Zāļu nosaukums:
Evalon (--) - Suspension and solvent for oral spray

Aktīvā viela:
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English

Mērķsugas:
Cālis

Lietošanas veids:
Iekšķīgai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:
Pieejamas tikai English
Presentation_strength:332-450 oocysts Reference:Hse Comments:Suspension Index:0
Pieejamas tikai English
Presentation_strength:196-265 oocysts Reference:Hse Comments:Suspension Index:1
Pieejamas tikai English
Presentation_strength:340-460 oocysts Reference:Hse Comments:Suspension Index:2
Pieejamas tikai English
Presentation_strength:213-288 oocysts Reference:Hse Comments:Suspension Index:3
Pieejamas tikai English
Presentation_strength:276-374 oocysts Reference:Hse Comments:Suspension Index:4

Farmaceitiskā forma:
Šī informācija par šīm zālēm nav pieejama.

Ierobežojumu periods pēc lietošanas veida:
Iekšķīgai lietošanai:

- **Cālis**
  - Visi attiecīgie audi. 0 diena Zero days

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QI01AN01

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

Iepakojuma apraksts:
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English

Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:
Pilns pieteikums - zināmai aktīvai vielai (Direktīvas 2001/82/EC 12.(3) pants)

Tirdzniecības atļaujas turētājs:
Laboratorios Hipra S.A.

Tirdzniecības atļaujas datums:
18/04/2016

Ražošanas vietas sēriju izlaidei:
Laboratorios Hipra S.A.

Atbildīgā iestāde:
European Commission

Atļaujas numurs:
Šī informācija par šīm zālēm nav pieejama.

Atļaujas statusa maiņas datums:
18/04/2016

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

Dokumenti

Visu dokumentu apvienotais fails

latviešu (PDF)
Publicēts: 17/06/2024
Lejupielādēt

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