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Syncroprost, 0.250 mg/ml solution for injection for cattle, horses, pigs and goats

Atļautas

- Cloprostenol

Zāļu identifikācija

Zāļu nosaukums:
Syncroprost, 0.250 mg/ml solution for injection for cattle, horses, pigs and goats

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Liellops (govs)
Liellops (tele)
Zirgs (ķēve)
Cūka (sieviešu kārtas)
Kaza (pieaugusi kaza)

Lietošanas veids:
intramuskulārai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

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

Farmaceitiskā forma:

Šķīdums injekcijām

Ierobežojumu periods pēc lietošanas veida:

intramuskulārai lietošanai:

- **Liellops (govs)**
  - Gaļa un blakusprodukti. 1 diena
  - Milk. 0 diena
- **Liellops (tele)**
  - Gaļa un blakusprodukti. 1 diena
  - Milk. 0 diena
- **Zirgs (ķēve)**
  - Gaļa un blakusprodukti. 1 diena
  - Milk. 0 diena
- **Cūka (sieviešu kārtas)**
  - Gaļa un blakusprodukti. 1 diena
- **Kaza (pieaugusi kaza)**
  - Gaļa un blakusprodukti. 1 diena
  - Milk. 0 diena

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:

QG02AD90

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

Pieejams:
Sweden

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:
Ceva Sante Animale

Tirdzniecības atļaujas datums:
17/07/2022

Ražošanas vietas sēriju izlaidei:
Vetem S.p.A.

Atbildīgā iestāde:
Swedish Medical Products Agency

Atļaujas numurs:
61562

Atļaujas statusa maiņas datums:

17/07/2022

Atsauces dalībvalsts:

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

Procedūras numurs:

IT/V/0147/001

Attiecīgās dalībvalstis:

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

Lietošanas instrukcija

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

Marķējuma teksts

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

Zāļu apraksts

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