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Procamidor Duo 40 mg/ml + 0,036 mg/ml Injektionslösung für Tiere

Atļautas

- Procaine hydrochloride
- Adrenaline hydrogen tartrate

Zāļu identifikācija

Zāļu nosaukums:
Procamidor Duo 40 mg/ml + 0,036 mg/ml Injektionslösung für Tiere

Aktīvā viela:
Pieejamas tikai English
Pieejamas tikai English

Mērķsugas:
Liellops
Aita
Zirgs
Cūka

Lietošanas veids:
perineirālai lietošanai
subkutānai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

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

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

Farmaceitiskā forma:

Šķīdums injekcijām

Ierobežojumu periods pēc lietošanas veida:

perineirālai lietošanai:

- **Liellops**
  - Milk. 0 stunda
  - Gaļa un blakusprodukti. 0 diena
- **Aita**
  - Milk. 0 stunda
  - Gaļa un blakusprodukti. 0 diena
- **Zirgs**
  - Milk. 0 stunda
  - Gaļa un blakusprodukti. 0 diena
- **Cūka**
  - Gaļa un blakusprodukti. 0 diena

subkutānai lietošanai:

- **Liellops**
  - Milk. 0 stunda
  - Gaļa un blakusprodukti. 0 diena
- **Aita**

- Milk. 0 stunda
- Gaļa un blakusprodukti. 0 diena

- **Zirgs**
  - Milk. 0 stunda
  - Gaļa un blakusprodukti. 0 diena

- **Cūka**
  - Gaļa un blakusprodukti. 0 diena

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QN01BA52

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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Pieejams:
Sweden

Iepakojuma apraksts:
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:
Vetviva Richter GmbH

Tirdzniecības atļaujas datums:
25/07/2019

Ražošanas vietas sēriju izlaidei:
Vetviva Richter GmbH

Atbildīgā iestāde:
Swedish Medical Products Agency

Atļaujas numurs:
57723

Atļaujas statusa maiņas datums:
25/07/2019

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

Procedūras numurs:
AT/V/0018/001

Attiecīgās dalībvalstis:
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Čehija
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Pieejamas tikai Spanish German Estonian English French Italian Dutch Portuguese Slovak Finnish Swedish Icelandic Norwegian

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Pieejamas tikai Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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

Dokumenti

Lietošanas instrukcija

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.

Marķējuma teksts

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

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