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Versiguard Rabies, Suspension for injection

Atļautas

- Rabies virus, strain SAD Vnukovo-32, Inactivated

Zāļu identifikācija

Zāļu nosaukums:
Versiguard Rabies, Suspension for injection

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Kaķis
Liellops
Cūka
Aita
Kaza
Zirgs
Mājas sesks
Suns

Lietošanas veids:
intramuskulārai lietošanai
subkutānai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

Pieejamas tikai English
2.00 starptautiskā(s) vienība(s) / 1.00 Deva

Farmaceitiskā forma:

Suspensija injekcijām

Ierobežojumu periods pēc lietošanas veida:
intramuskulārai lietošanai:

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

subkutānai lietošanai:

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

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QI07AA02

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:

Norway

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

Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:
Pieejamas tikai English Italian

Tirdzniecības atļaujas turētājs:
Zoetis Animal Health ApS

Tirdzniecības atļaujas datums:
22/05/2009

Ražošanas vietas sēriju izlaidei:
Bioveta a.s.

Atbildīgā iestāde:
Norwegian Medical Products Agency

Atļaujas numurs:
05-3851

Atļaujas statusa maiņas datums:
29/03/2011

Atsauces dalībvalsts:
Čehija

Procedūras numurs:
CZ/V/0100/001

Attiecīgās dalībvalstis:

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

Dokumenti

Zāļu apraksts

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.