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Bovilis Rotavec Corona Emulsion for Injection for Cattle

Atļautas

- Escherichia coli, serotype O101:K99 (fimbrial adhesins F5 and F41), strain CN7985, Inactivated
- Bovine rotavirus A, type G6P5, strain UK-Compton, Inactivated
- Bovine coronavirus, strain Mebus, Inactivated

Zāļu identifikācija

Zāļu nosaukums:
Bovilis Rotavec Corona Emulsion for Injection for Cattle

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

Mērķsugas:
Liellops

Lietošanas veids:
intramuskulārai lietošanai
subkutānai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

Pieejamas tikai English
560.00 Cietfāzes enzīmu imūnosorbences testa vienība(s) / 2.00 mililitrs(i)

Pieejamas tikai English
874.00 Cietfāzes enzīmu imūnosorbences testa vienība(s) / 2.00 mililitrs(i)

Pieejamas tikai English
340.00 Cietfāzes enzīmu imūnosorbences testa vienība(s) / 2.00 mililitrs(i)

Farmaceitiskā forma:

Emulsija injekcijām

Ierobežojumu periods pēc lietošanas veida:

intramuskulārai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 0 diena
  - Piens. 0 diena

subkutānai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 0 diena
  - Piens. 0 diena

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:

QI02AL01

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:
Denmark

Iepakojuma apraksts:
Pieejamas tikai English
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 imunoloģiskām veterinārajām zālēm (Direktīvas 2001/82/EC 13.(d) pants)

Tirdzniecības atļaujas turētājs:
Intervet International B.V.

Tirdzniecības atļaujas datums:
21/02/2008

Ražošanas vietas sēriju izlaidei:
Intervet International B.V.

Atbildīgā iestāde:
Danish Medicines Agency

Atļaujas numurs:
41975

Atļaujas statusa maiņas datums:
21/02/2008

Atsauces dalībvalsts:

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

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Procedūras numurs:

DE/V/0276/001

Attiecīgās dalībvalstis:

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

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Pieejamas tikai Spanish Czech German Estonian Greek 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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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.

Visu dokumentu apvienotais fails

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