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Nobilis RT+IBmulti+ND+EDS emulsion for injection for chickens

Atļautas

- Newcastle disease virus, strain Clone 30, Inactivated
- Eggdrop syndrome-1976 virus, strain BC14, Inactivated
- Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
- Infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Zāļu identifikācija

Zāļu nosaukums:
Nobilis RT+IBmulti+ND+EDS emulsion for injection for chickens

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

Mērķsugas:
Cālis

Lietošanas veids:
intramuskulārai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:

Pieejamas tikai English
50.00 50% aizsardzības deva / 0.50 mililitrs(i)

Pieejamas tikai English
6.50 log2 hemaglutinācijas inhibēšanas vienība(s) / 0.50 mililitrs(i)

Pieejamas tikai English
9.50 log2 cietfāzes enzīmu imūnosorbences testa vienība(s) / 0.50 mililitrs(i)

Pieejamas tikai English
4.00 log2 vīrusu neitralizēšanas vienība(s) / 0.50 mililitrs(i)

Pieejamas tikai English
5.50 log2 vīrusu neitralizēšanas vienība(s) / 0.50 mililitrs(i)

Farmaceitiskā forma:

Emulsija injekcijām

Ierobežojumu periods pēc lietošanas veida:

intramuskulārai lietošanai:

- **Cālis**
  - Gaļa un blakusprodukti. 0 diena
  - Ola. 0 diena

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:

QI01AA18

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:
Ireland

Iepakojuma apraksts:
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 (Ireland) Limited

Tirdzniecības atļaujas datums:
3/11/2004

Ražošanas vietas sēriju izlaidei:
Intervet International B.V.

Atbildīgā iestāde:
Health Products Regulatory Authority

Atļaujas numurs:
VPA10996/181/001

Atļaujas statusa maiņas datums:
3/11/2004

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

Procedūras numurs:
DE/V/0209/001

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 English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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

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

Visu dokumentu apvienotais fails

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