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CANIGEN CHPPI/L LYOPHILISATE AND SUSPENSION FOR SUSPENSION FOR INJECTION FOR DOGS

Atļautas

- Canine adenovirus 2, strain Manhattan, Live
- Canine parainfluenza virus, strain Manhattan, Live
- Canine parvovirus, strain Cornell 780916, Live
- Leptospira interrogans, serovar Icterohaemorrhagiae, strain 601895, Inactivated
- Leptospira interrogans, serovar Canicola, strain 601903, Inactivated
- Canine distemper virus, strain Lederle, Live

Zāļu identifikācija

Zāļu nosaukums:
CANIGEN CHPPI/L LYOPHILISATE AND SUSPENSION FOR SUSPENSION FOR INJECTION FOR DOGS

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

Mērķsugas:
Suns

Lietošanas veids:
subkutānai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:
Pieejamas tikai English
10000.00 šūnu kultūras infekciozā deva 50 / 1.00 Deva

Pieejamas tikai English
100000.00 šūnu kultūras infekciozā deva 50 / 1.00 Deva

Pieejamas tikai English
100000.00 šūnu kultūras infekciozā deva 50 / 1.00 Deva

Pieejamas tikai English
4250.00 Cietfāzes enzīmu imūnosorbences testa vienība(s) / 1.00 Deva

Pieejamas tikai English
4350.00 Cietfāzes enzīmu imūnosorbences testa vienība(s) / 1.00 Deva

Pieejamas tikai English
1000.00 šūnu kultūras infekciozā deva 50 / 1.00 Deva

Farmaceitiskā forma:
Liofilizāts un suspensija suspensijas injekcijām pagatavošanai

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QI07AI02

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:
Poland

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:
Pilns pieteikums - zināmai aktīvai vielai (Direktīvas 2001/82/EC 12.(3) pants)

Tirdzniecības atļaujas turētājs:
Virbac

Tirdzniecības atļaujas datums:
22/11/2016

Ražošanas vietas sēriju izlaidei:
Virbac

Atbildīgā iestāde:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Atļaujas numurs:
2585

Atļaujas statusa maiņas datums:
22/11/2016

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

Procedūras numurs:

FR/V/0237/001

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 Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Pieejamas tikai Spanish Czech German Estonian English French Croatian 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

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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.

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.

Visu dokumentu apvienotais fails

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

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