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

Ima dovoljenje za promet

- 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

## Informacije o zdravilu

**Ime zdravila:**
CANIGEN CHPPI/L LYOPHILISATE AND SUSPENSION FOR SUSPENSION FOR INJECTION FOR DOGS

**Učinkovina:**
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

**Ciljne živalske vrste:**
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

**Pot uporabe:**
Subkutana uporaba

## Podatki o zdravilu

**Učinkovina / Jakost:**
Na voljo samo v English
10000.00 50% cell culture infectious dose / 1.00 Dose

Na voljo samo v English
100000.00 50% cell culture infectious dose / 1.00 Dose

Na voljo samo v English
100000.00 50% cell culture infectious dose / 1.00 Dose

Na voljo samo v English
4250.00 enzyme-linked immunosorbent assay unit / 1.00 Dose

Na voljo samo v English
4350.00 enzyme-linked immunosorbent assay unit / 1.00 Dose

Na voljo samo v English
1000.00 50% cell culture infectious dose / 1.00 Dose

**Farmacevtska oblika:**
Liofilizat in suspenzija za suspenzijo za injiciranje

**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QI07AI02

**Pravni status za dobavo/izdajo zdravila:**
Zdravilo, ki se izdaja na veterinarski recept

**Status dovoljenja za promet z zdravilom:**
Valid

**Ima dovoljenje za promet v:**

Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

**Na voljo v:**
Greece

**Opis ovojnine zdravila:**
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English

## Dodatne informacije

**Vrsta dovoljenja:**
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

**Pravna podlaga za izdajo dovoljenja za promet z zdravilom:**
Na voljo samo v English Italian Latvian Lithuanian Norwegian

**Imetnik dovoljenja za promet z zdravilom:**
Virbac

**Datum dovoljenja za promet z zdravilom:**
6/03/2016

**Proizvodna mesta, odgovorna za sproščanje serij:**
Virbac

**Pristojni organ:**
National Organization For Medicines

**Številka dovoljenja za promet z zdravilom:**
105536/27-10-2020/K-0217301

**Datum spremembe statusa dovoljenja za promet:**
26/10/2020

**Referenčna država članica:**
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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**Številka postopka:**
FR/V/0237/001

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**Zadevne države članice:**
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

## Dokumenti

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

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