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Biocan Novel Pi L4, Lyophilisate and solvent for suspension for injection

Ima dovoljenje za promet

- Canine parainfluenza virus, strain CPiV-2-Bio 15, Live
- Leptospira interrogans, serovar Icterohaemorrhagiae, strain MSLB 1089, Inactivated
- Leptospira interrogans, serovar Canicola, strain MSLB 1090, Inactivated
- Leptospira interrogans, serovar Bratislava, strain MSLB 1088, Inactivated
- Leptospira kirschneri, serovar Grippotyphosa, strain MSLB 1091, Inactivated

Informacije o zdravilu

Ime zdravila:
Biocan Novel Pi L4, Lyophilisate and solvent for suspension for injection

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

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
5.10 log10 50% tissue culture infectious dose / 1.00 Dose

Na voljo samo v English
51.00 Antibody microagglutination-lytic reaction / 1.00 Dose

Na voljo samo v English
51.00 Antibody microagglutination-lytic reaction / 1.00 Dose

Na voljo samo v English
51.00 Antibody microagglutination-lytic reaction / 1.00 Dose

Na voljo samo v English
40.00 Antibody microagglutination-lytic reaction / 1.00 Dose

Farmacevtska oblika:
Liofilizat in vehikel za suspenzijo za injiciranje

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI07AI08

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

Opis ovojnine zdravila:

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

Imetnik dovoljenja za promet z zdravilom:
Bioveta a.s.

Datum dovoljenja za promet z zdravilom:
2/10/2014

Proizvodna mesta, odgovorna za sproščanje serij:
Bioveta a.s.

Pristojni organ:
Food And Veterinary Service

Številka dovoljenja za promet z zdravilom:
V/DCP/14/0051

Datum spremembe statusa dovoljenja za promet:
2/10/2014

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

Številka postopka:
CZ/V/0123/001

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 Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian Greek 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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