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Nobivac L4 (--) - Suspension for injection

Ima dovoljenje za promet

- Leptospira interrogans, serovar Copenhageni, strain Ic-02-001, Inactivated
- Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated
- Leptospira interrogans, serovar Bratislava, strain As-05-073, Inactivated
- Leptospira kirschneri, serovar Dadas, strain GR-01-005, Inactivated

Informacije o zdravilu

Ime zdravila:
Nobivac L4 (--) - Suspension for injection

Učinkovina:
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
Presentation_strength:≥290 U Reference:Hse Index:0
Na voljo samo v English
Presentation_strength:≥3550 U Reference:Hse Index:1
Na voljo samo v English
Presentation_strength:≥500 U Reference:Hse Index:2
Na voljo samo v English
Presentation_strength:≥650 U Reference:Hse Index:3

Farmacevtska oblika:
Suspenzija za injiciranje

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI07AB01

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

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Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Na voljo v:

Austria , Belgium , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Portugal , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

Opis ovojnine zdravila:

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

Imetnik dovoljenja za promet z zdravilom:
Intervet International B.V.

Datum dovoljenja za promet z zdravilom:
16/07/2012

Proizvodna mesta, odgovorna za sproščanje serij:
Intervet International B.V.

Pristojni organ:
European Commission

Številka dovoljenja za promet z zdravilom:
Ta podatek za to zdravilo ni na voljo.

Datum spremembe statusa dovoljenja za promet:
16/07/2012

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Combined File of all Documents

slovenščina (PDF)
Objavljeno na dan: 13/03/2026
Prenesi

ema-puar-nobivac-l4-v-2010-par-en.pdf

ema-puar-v2010-nobivacl4-ws2673-vra0000184065-en.pdf