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Versican Plus L4 (--) - Suspension for injection

Ima dovoljenje za promet

- Leptospira interrogans, serovar Icterohaemorrhagiae, strain MSLB 1089, Inactivated
- Leptospira interrogans, serovar Canicola, strain MSLB 1090, Inactivated
- Leptospira kirschneri, serovar Grippotyphosa, strain MSLB 1091, Inactivated
- Leptospira interrogans, serovar Bratislava, strain MSLB 1088, Inactivated

Informacije o zdravilu

Ime zdravila:
Versican Plus 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
1.51 Antibody microagglutination-lytic reaction / 1.00 Viala

Na voljo samo v English
1.51 Antibody microagglutination-lytic reaction / 1.00 Viala

Na voljo samo v English
1.40 Antibody microagglutination-lytic reaction / 1.00 Viala

Na voljo samo v English
Presentation_strength:1:51 ALR Reference:Ph EUR 0447 Comments:Suspension 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 Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Na voljo v:
Austria , Belgium , Cyprus , Denmark , Germany , Greece , Ireland , Italy , Luxembourg , Netherlands , United Kingdom (Northern Ireland)

Opis ovojnine zdravila:

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 Norwegian

Imetnik dovoljenja za promet z zdravilom:
Zoetis Belgium

Datum dovoljenja za promet z zdravilom:
31/07/2014

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

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
31/07/2014

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: 7/09/2026
Prenesi

ema-puar-versican-plus-l4-v-3680-par-en.pdf