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Leucofeligen FeLV/RCP (--) - Lyophilisate and suspension for suspension for injection

Ima dovoljenje za promet

- Feline panleucopenia virus, strain LR 72, Live
- Feline calicivirus, strain F9, Live
- Felid herpesvirus 1, strain F2, Live
- Feline leukemia virus, envelope protein p45

Informacije o zdravilu

Ime zdravila:
Leucofeligen FeLV/RCP (--) - Lyophilisate and suspension 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

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:$10^{3.7}$-$10^{4.5}$ $CCID_{50}$ Reference:Hse Comments:Freeze dried component Index:0
Na voljo samo v English
Presentation_strength:$10^{4.6}$-$10^{6.1}$ $CCID_{50}$ Reference:Hse Comments:Freeze dried component Index:1
Na voljo samo v English
Presentation_strength:$10^{5.0}$-$10^{6.6}$ $CCID_{50}$ Reference:Hse Comments:Freeze dried component Index:2
Na voljo samo v English
102.00 microgram(s) / 1.00 Viala

Farmacevtska oblika:
Liofilizat in suspenzija za suspenzijo za injiciranje

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI06AH07

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 , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Poland , Portugal , Romania , Slovakia , Slovenia , 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

Imetnik dovoljenja za promet z zdravilom:
Virbac

Datum dovoljenja za promet z zdravilom:
25/06/2009

Proizvodna mesta, odgovorna za sproščanje serij:
Virbac

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
25/06/2009

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: 19/10/2026
Prenesi

ema-puar-leucofeligen-v-143-par-en.pdf

ema-puar-leucofeligen-var-ws1282-en.pdf