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

Atļautas

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

Zāļu identifikācija

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

Aktīvā viela:
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English
Pieejamas tikai English

Mērķsugas:
Kaķis

Lietošanas veids:
subkutānai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:
Pieejamas tikai English
Presentation_strength:$10^{3.7}$-$10^{4.5}$ $CCID_{50}$ Reference:Hse Comments:Freeze dried component Index:0
Pieejamas tikai English
Presentation_strength:$10^{4.6}$-$10^{6.1}$ $CCID_{50}$ Reference:Hse Comments:Freeze dried component Index:1
Pieejamas tikai English
Presentation_strength:$10^{5.0}$-$10^{6.6}$ $CCID_{50}$ Reference:Hse Comments:Freeze dried component Index:2
Pieejamas tikai English
102.00 mikrograms (i) / 1.00 Flakons

Farmaceitiskā forma:
Liofilizāts un suspensija suspensijas injekcijām pagatavošanai

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Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QI06AH07

Izplatīšanas juridiskie nosacījumi:
Recepšu veterinārās zāles

Atļaujas statuss:
Derīga

Atļautas:
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Pieejamas tikai Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Pieejams:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Denmark , Estonia , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Poland , Portugal , Romania , Slovakia , Slovenia , United Kingdom (Northern Ireland) , Čehija

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Iepakojuma apraksts:

Pieejamas tikai English

Pieejamas tikai English

Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:
Pieejamas tikai English Italian

Tirdzniecības atļaujas turētājs:
Virbac

Tirdzniecības atļaujas datums:
25/06/2009

Ražošanas vietas sēriju izlaidei:
Virbac

Atbildīgā iestāde:
European Commission

Atļaujas numurs:
Šī informācija par šīm zālēm nav pieejama.

Atļaujas statusa maiņas datums:
25/06/2009

Lai uzzinātu par veterināro zāļu blakusparādībām, lūdzu, dodieties uz: www.adrreports.eu/vet

Dokumenti

Visu dokumentu apvienotais fails

latviešu (PDF)
Publicēts: 19/10/2026

Lejupielādēt

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

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