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Purevax RCPCh (--) - Lyophilisate and solvent for suspension for injection

Ima dovoljenje za promet

- Feline calicivirus, strains 431 and G1, Inactivated
- Chlamydia felis, strain 905, Live
- Feline panleucopenia virus, strain PLI IV, Live
- Felid herpesvirus 1, strain F2, Live

Informacije o zdravilu

Ime zdravila:
Purevax RCPCh (--) - 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

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:≥2.0 ELISA U Reference:Hse Index:0
Na voljo samo v English
Presentation_strength:≥$10^{3.0}$ EID_{50} Reference:Hse Index:1
Na voljo samo v English
Presentation_strength:≥$10^{3.5}$ $CCID_{50}$ Reference:Hse Index:2
Na voljo samo v English
Presentation_strength:≥$10^{4.9}$ $CCID_{50}$ Reference:Hse Index:3

Farmacevtska oblika:
Liofilizat in vehikel za suspenzijo za injiciranje

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI06AX

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 v:
Austria , Belgium , Czechia , France , Luxembourg , Poland , Slovakia , Spain

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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

Imetnik dovoljenja za promet z zdravilom:
Boehringer Ingelheim Vetmedica GmbH

Datum dovoljenja za promet z zdravilom:
23/02/2005

Proizvodna mesta, odgovorna za sproščanje serij:
Boehringer Ingelheim Animal Health France

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
23/02/2005

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: 14/03/2026

Prenesi

ema-puar-purevax-rcpch-v-088-par-en.pdf