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ProteqFlu (--) - Suspension for injection

Ima dovoljenje za promet

- Canarypox virus, strain vCP3011, expressing haemagglutinin gene of Influenza A virus subtype H3N8 (A/equine/Richmond/1/2007), Live
- Canarypox virus, strain vCP2242, expressing haemagglutinin gene of Influenza A virus subtype H3N8 (A/equine/Ohio/03), Live

Informacije o zdravilu

Ime zdravila:
ProteqFlu (--) - Suspension for injection

Učinkovina:
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:
intramuskularna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
Presentation_strength:vCP3011 ≥ $10^{5,3}$ FAID50 Reference:Hse Index:0
Na voljo samo v English
Presentation_strength:vCP2242 ≥ $10^{5,3}$ FAID50 Reference:Hse Index:1

Farmacevtska oblika:
Suspenzija za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **Horse**
  - Not applicable. 0 day Zero days

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI05AD02

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:
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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 , Czechia , France , Luxembourg , Poland , Slovakia , Spain

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Opis ovojnine zdravila:

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:
6/03/2003

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:
13/02/2006

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-proteqflu-v-073-var-ii-0014-en.pdf

ema-puar-proteqflu-v-073-par-en.pdf