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Porcilis PCV M Hyo ID (--) + (--) - Emulsion for injection

Ima dovoljenje za promet

- Porcine circovirus type 2, ORF2 capsid protein
- Mycoplasma hyopneumoniae, strain J, Inactivated

Informacije o zdravilu

Ime zdravila:
Porcilis PCV M Hyo ID (--) + (--) - Emulsion 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:
Intradermalna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
Presentation_strength:751.4 AU Reference:HSE Index:0

Na voljo samo v English
Presentation_strength:0.72 AU Reference:HSE Index:1

Farmacevtska oblika:
Emulzija za injiciranje

Karenca glede na pot uporabe:
Intradermalna uporaba:

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

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QI09AL08

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 v:
Portugal

Opis ovojnine zdravila:
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
Na voljo samo v English
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 French Italian Latvian Lithuanian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Intervet International B.V.

Datum dovoljenja za promet z zdravilom:
30/08/2024

Proizvodna mesta, odgovorna za sproščanje serij:
Intervet International B.V.

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
30/08/2024

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-v6289-porcilispcvmhyoid-en.pdf