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Zuprevo 40 mg/ml - Solution for injection

Ima dovoljenje za promet

- Tildipirosin

Informacije o zdravilu

Ime zdravila:
Zuprevo 40 mg/ml - Solution for injection

Učinkovina:
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
40.00 milligram(s) / 1.00 Viala

Farmacevtska oblika:

Raztopina za injiciranje

Karenca glede na pot uporabe:

intramuskularna uporaba:

- **Pig**
  - Meat and offal. 9 day 9 days

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

QJ01FA96

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 Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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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 , Czechia , Denmark , Germany , Hungary , Italy , 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

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English Italian

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

Datum dovoljenja za promet z zdravilom:
6/05/2011

Proizvodna mesta, odgovorna za sproščanje serij:
Intervet International GmbH

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
6/05/2011

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-zuprevo-v-2009-var-ii-0006-g-en.pdf

ema-puar-zuprevo-v-2009-par-en.pdf