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Naxcel 200 mg/ml - Suspension for injection

Ima dovoljenje za promet

- Ceftiofur

Informacije o zdravilu

Ime zdravila:
Naxcel 200 mg/ml - Suspension for injection

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
govedo

Pot uporabe:
Subkutana uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
200.00 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Suspenzija za injiciranje

Karenca glede na pot uporabe:

Subkutana uporaba:

- **govedo**
  - Meat and offal. 9 day 9 days
  - Milk. 0 day Zero days

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ01DD90

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 Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Na voljo v:
Austria , Belgium , Cyprus , Czechia , Estonia , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , 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

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Pravna podlaga za izdajo dovoljenja za promet z zdravilom:

Na voljo samo v English

Imetnik dovoljenja za promet z zdravilom:

Zoetis Belgium

Datum dovoljenja za promet z zdravilom:
19/05/2005

Proizvodna mesta, odgovorna za sproščanje serij:
Zoetis Belgium

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
8/10/2009

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-naxcel-v-079-var-ii-0012-en.pdf

ema-puar-naxcel-v-079-par-en.pdf