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Naxcel 200 mg/ml - Suspension for injection

Atļautas

- Ceftiofur

Zāļu identifikācija

Zāļu nosaukums:
Naxcel 200 mg/ml - Suspension for injection

Aktīvā viela:
Pieejamas tikai English

Mērķsugas:
Liellops

Lietošanas veids:
subkutānai lietošanai

Sīkāka informācija par zālēm

Aktīvā viela un stiprums:
Pieejamas tikai English
200.00 miligrams(i) / 1.00 mililitrs(i)

Farmaceitiskā forma:
Suspensija injekcijām

Ierobežojumu periods pēc lietošanas veida:

subkutānai lietošanai:

- **Liellops**
  - Gaļa un blakusprodukti. 9 diena 9 days
  - Piens. 0 diena Zero days

Anatomiski terapeitiski ķīmiskais veterinārais (ATĶvet) kods:
QJ01DD90

Izplatīšanas juridiskie nosacījumi:
Recepšu veterinārās zāles

Atļaujas statuss:
Derīga

Atļautas:
Pieejamas tikai Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Pieejamas tikai Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

Pieejams:
Austria , Belgium , Cyprus , Estonia , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Poland , Slovakia , Spain , Čehija

Iepakojuma apraksts:
Pieejamas tikai English

Papildu informācija

Tiesību tips:

Veterināro zāļu atļaujas piešķiršanas juridiskais pamats:
Pieejamas tikai English

Tirdzniecības atļaujas turētājs:
Zoetis Belgium

Tirdzniecības atļaujas datums:
19/05/2005

Ražošanas vietas sēriju izlaidei:
Zoetis Belgium

Atbildīgā iestāde:
European Commission

Atļaujas numurs:
Šī informācija par šīm zālēm nav pieejama.

Atļaujas statusa maiņas datums:
8/10/2009

Lai uzzinātu par veterināro zāļu blakusparādībām, lūdzu, dodieties uz: www.adrreports.eu/vet

Dokumenti

Visu dokumentu apvienotais fails

latviešu (PDF)
Publicēts: 14/03/2026
Lejupielādēt

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

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