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Lexylan 180 mg/ml - Suspension for injection

Ima dovoljenje za promet

- Cefalexin

Informacije o zdravilu

Ime zdravila:
Lexylan 180 mg/ml - Suspension for injection

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
govedo
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
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
Subkutana uporaba

Podatki o zdravilu

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

Farmacevtska oblika:
Suspenzija za injiciranje

Karenca glede na pot uporabe:
intramuskularna uporaba:

- **govedo**
  - Meat and offal. 12 day — 12 Days
  - Milk. 0 hour — Zero Hours

Subkutana uporaba:

- **govedo**
  - Meat and offal. 12 day — 12 Days
  - Milk. 0 hour — Zero Hours

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ01DB01

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

Na voljo v:
Austria , Belgium , Denmark , France , Germany , Ireland , Italy , Luxembourg , Netherlands , Spain

Opis ovojnine zdravila:
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:
Emdoka

Datum dovoljenja za promet z zdravilom:
8/04/2024

Proizvodna mesta, odgovorna za sproščanje serij:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Pristojni organ:
European Commission

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

Datum spremembe statusa dovoljenja za promet:
8/04/2024

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: 13/03/2026
Prenesi

ema-puar-v6103-lexylan-en.pdf