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# LABIMYCIN LA 300 mg/ml SOLUTION FOR INJECTION

Ima dovoljenje za promet

- Oxytetracycline dihydrate

## Informacije o zdravilu

**Ime zdravila:**
LABIMYCIN LA 300 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
govedo
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
323.50 milligram(s) / 1.00 millilitre(s)

**Farmacevtska oblika:**

Raztopina za injiciranje

**Karenca glede na pot uporabe:**

**intramuskularna uporaba:**

- **Pig**
  - Meat and offal. 28 day
- **govedo**
  - Meat and offal. 35 day
  - Milk. 10 day
- **Sheep**
  - Meat and offal. 35 day
  - Milk. 9 day

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

QJ01AA06

**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 Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

**Na voljo v:**

Spain

**Opis ovojnine zdravila:**
Na voljo samo v Spanish
Na voljo samo v Spanish
Na voljo samo v Spanish
Na voljo samo v Spanish
Na voljo samo v Spanish
Na voljo samo v Spanish

## 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 Italian Latvian Norwegian

**Imetnik dovoljenja za promet z zdravilom:**
Labiana Life Sciences S.A.

**Datum dovoljenja za promet z zdravilom:**
5/04/2022

**Proizvodna mesta, odgovorna za sproščanje serij:**
Labiana Life Sciences S.A.

**Pristojni organ:**
Spanish Agency Of Medicines And Medical Devices

**Številka dovoljenja za promet z zdravilom:**
4074 ESP

**Datum spremembe statusa dovoljenja za promet:**
6/04/2022

**Referenčna država članica:**
Na voljo samo v Spanish Czech German Estonian English French Croatian Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

**Številka postopka:**
ES/V/0406/001

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**Zadevne države članice:**
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

## Dokumenti

Navodilo za uporabo

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Označevanje

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Povzetek glavnih značilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

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