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Orbenin LA 200 mg intramammary suspension for lactating cattle and sheep

Ima dovoljenje za promet

- Cloxacillin sodium monohydrate

Informacije o zdravilu

Ime zdravila:
Orbenin LA 200 mg intramammary suspension for lactating cattle and sheep

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
Na voljo samo v Spanish Czech Danish Estonian Greek English French Italian Latvian Lithuanian Romanian Finnish Swedish 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:
Intramamarno dajanje

Podatki o zdravilu

Učinkovina / Jakost:

Na voljo samo v English
210.08 milligram(s) / 1.00 Aplikator

Farmacevtska oblika:

intramamarna suspenzija

Karenca glede na pot uporabe:

Intramamarno dajanje:

- **Cattle (dairy cow)**
  - Milk. 96 hour
  - Meat and offal. 7 day
- **Sheep**
  - Meat and offal. 7 day
  - Milk. no withdrawal period

Not authorised for use in sheep producing milk for human consumption

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

QJ51CF02

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

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

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

Imetnik dovoljenja za promet z zdravilom:
Zoetis Portugal Lda.

Datum dovoljenja za promet z zdravilom:
8/11/2019

Proizvodna mesta, odgovorna za sproščanje serij:
Haupt Pharma Latina S.r.l.

Pristojni organ:
Directorate General For Food And Veterinary

Številka dovoljenja za promet z zdravilom:
1312/01/19RFVPT

Datum spremembe statusa dovoljenja za promet:
17/10/2022

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

Številka postopka:
DE/V/0319/001

Zadevne države članice:

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

Combined File of all Documents

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

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