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Synulox LC Plus Intramammary suspension for lactating cattle

Ima dovoljenje za promet

- Prednisolone
- Potassium clavulanate
- Amoxicillin trihydrate

Informacije o zdravilu

Ime zdravila:
Synulox LC Plus Intramammary suspension for lactating cattle

Učinkovina:
Na voljo samo v English
Na voljo samo v English
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

Pot uporabe:
Intramamarno dajanje

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English

10.00 milligram(s) / 1.00 Aplikator

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

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

Farmacevtska oblika:
intramamarna suspenzija

Karenca glede na pot uporabe:
Intramamarno dajanje:

- **Cattle (dairy cow)**
  - Meat and offal. 7 day
  - Milk. 84 hour

84 hours. With cows milked twice daily, milk for human consumption may only be taken the 7th milking after the last treatment. Where any other milking routine is followed, milk may be taken for human consumption only after the same period from the last treatment (e.g. with 3 times a day milking, milk may be taken for human consumption at the 11th milking).

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ51RV01

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 v:
Germany

Opis ovojnine zdravila:
Na voljo samo v English
Na voljo samo v English
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 Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Zoetis Deutschland GmbH

Datum dovoljenja za promet z zdravilom:
29/10/2013

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

Pristojni organ:
Federal Office Of Consumer Protection And Food Safety

Številka dovoljenja za promet z zdravilom:
401895.00.00

Datum spremembe statusa dovoljenja za promet:
19/12/2018

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

Številka postopka:
DE/V/0315/001

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 Estonian English French Lithuanian Portuguese 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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