

LINEOMAM LC, Intramammary solution

Authorised

- Lincomycin
- NEOMYCIN SULFATE

Product identification

Medicine name:

LINEOMAM LC, Intramammary solution

Lineomam LC 330 mg/10 ml + 100 000 IU/10 ml šķīdums ievadīšanai tesmenī

Active substance:

Lincomycin

NEOMYCIN SULFATE

Target species:

Cattle (lactating cow)

Route of administration:

Intramammary use

Product details

Active substance and strength:

Lincomycin

330.00 milligram(s) / 1.00 Applicator

NEOMYCIN SULFATE

100000.00 international unit(s) / 1.00 Applicator

Pharmaceutical form:

Intramammary solution

Withdrawal period by route of administration:**Intramammary use:**

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Cattle (lactating cow)

- Meat and offal. 3 day
 - Milk. 84 hour
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RF03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

Package description:

Plastic Applicator 24 x 1.0 Applicator

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Bioveta a.s.

Marketing authorisation date:

31/05/2017

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/MRP/17/0024

Date of authorisation status change:

31/05/2017

Reference member state:

Czechia

Procedure number:

CZ/V/0138/001

Concerned member states:

Bulgaria Croatia Cyprus Estonia Greece Hungary Latvia Lithuania Poland
Romania Slovakia Slovenia

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Package Leaflet

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Labelling

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Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

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