LINEOMAM LC, Intramammary solution

Authorised

- Lincomycin
- NEOMYCIN SULFATE

Product identification

Medicine name:
LINEOMAM LC 330 mg/10 ml + 100,000 IU/10 ml ενδομαστικό διάλυμα
LINEOMAM LC, Intramammary solution

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:

- **Cattle (lactating cow)**
  - Meat and offal. 3 day
  - Milk. 84 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RF03

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

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:
7/12/2017

Manufacturing sites for batch release:
Bioveta, a.s.

Responsible authority:
National Organization For Medicines

Authorisation number:
72522/28-07-2021/K-0226401

Date of authorisation status change:
27/07/2021

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