PENTOMAST LC, intramaminė suspensija galvijams

Authorised

- Prednisolone
- NEOMYCIN SULFATE
- Streptomycin sulfate
- Benzylpenicillin procaine

Product identification

Medicine name:
PENTOMAST LC, intramaminė suspensija galvijams

Active substance:
Prednisolone

NEOMYCIN SULFATE

Streptomycin sulfate

Benzylpenicillin procaine

Target species:
Cattle (cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:

Prednisolone
10.00 milligram(s) / 1.00 Syringe

NEOMYCIN SULFATE
100.00 milligram(s) / 1.00 Syringe

Streptomycin sulfate
100.00 milligram(s) / 1.00 Syringe

Benzylpenicillin procaine
100000.00 international unit(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (cow)**
  - Milk. 108 hour

108 hours after last treatment. The milk is not authorized for human consumption during treatment.

  - Meat and offal. 7 day

Slaughter of animals during treatment is not authorized.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RV01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:
Lithuania

Package description:
Available only in Lithuanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Interchemie werken De Adelaar LT UAB

Marketing authorisation date:
11/02/2018

Manufacturing sites for batch release:
Interchemie Werken De Adelaar Eesti AS

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/18/2447/001

Date of authorisation status change:
23/11/2025

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

Documents

Combined File of all Documents

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