LINKOLAKTAN, 750 mg, intramaminis gelis melžiamoms karvėms

Authorised

- Lincomycin

Product identification

Medicine name:
LINKOLAKTAN, 750 mg, intramaminis gelis melžiamoms karvėms

Active substance:
Lincomycin

Target species:
Cattle (lactating cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Lincomycin
750.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary gel

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (lactating cow)**
  - Meat and offal. 3 day
  - Milk. 84 hour 84 Hours or 7 milkings.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51FF02

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

Authorisation status:
Valid

Authorised in:
Lithuania

Available in:
Lithuania

Package description:
Available only in Lithuanian
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:

Fatro S.p.A.

Marketing authorisation date:
27/07/2015

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/15/2308/001-002

Date of authorisation status change:
25/09/2025

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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