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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:

Withdrawal period by route of administration: Intramammary use:

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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 <u>Lithuanian</u>
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:



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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