Galactis 750 mg gel intramammario per bovine in lattazione

Authorised

- Lincomycin hydrochloride

Product identification

Medicine name:
Galactis 750 mg gel intramammario per bovine in lattazione

Active substance:
Lincomycin hydrochloride

Target species:
Cattle (lactating cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Lincomycin hydrochloride
850.00 milligram(s) / 5.00 gram(s)

Pharmaceutical form:

Intramammary gel

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:

QJ51FF

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Available in:

Italy

Package description:

Available only in Italian

Available only in Italian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Fatro S.p.A.

Marketing authorisation date:
3/01/2013

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

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
29/01/2013

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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