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LINCOMASTINA 750 mg SOLUCION INTRAMAMARIA

Authorised

• Lincomycin hydrochloride

Product identification

Medicine name:

LINCOMASTINA 750 mg SOLUCION INTRAMAMARIA

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) / 1.00 Syringe

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:

QJ51FF02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Package description:

Available only in Spanish Available only in Spanish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Fatro S.p.A.

Marketing authorisation date:

4/11/2014

Manufacturing sites for batch release: Fatro S.p.A.
Responsible authority: Spanish Agency For Medicines And Medical Devices
Authorisation number: 3132 ESP
Date of authorisation status change: 4/11/2014
To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet
Documents
Summary of Product Characteristics
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Package Leaflet
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Labelling

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