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

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

es-puar-lincomastina-es.pdf