Qivitan LC 75 mg intramammary ointment for lactating cows

Authorised

• Cefquinome sulfate

Product identification

Medicine name:

Qivitan LC 75 mg intramammary ointment for lactating cows
QIVITAN LACTACION 75mg POMADA INTRAMAMARIA PARA VACAS EN LACTACION

Active substance:

Cefquinome sulfate

Target species:

Cattle

Route of administration:

Intramammary use

Product details

Active substance and strength:

Cefquinome sulfate 88.92 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary ointment

Withdrawal period by route of administration:

Intramammary use:

Cattle

- Meat and offal. 4 day
- Milk. 120 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51DE90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Package description:

Prefilled 8 g single-dose intramammary syringe consisting of white opaque LDPEbarrel with white opaque LDPE plunger and white opaque LDPE cap. Cleaning towels (smooth, white crepe paper impregnated with isopropylalcohol/benzalkonium chloride) individually wrapped. Cardboard boxes of 36 syringes and 36 cleaning towels.

Prefilled 8 g single-dose intramammary syringe consisting of white opaque LDPEbarrel with white opaque LDPE plunger and white opaque LDPE cap. Cleaning towels (smooth, white crepe paper impregnated with isopropylalcohol/benzalkonium chloride) individually wrapped. Cardboard boxes of 24 syringes and 24 cleaning towels.

Prefilled 8 g single-dose intramammary syringe consisting of white opaque LDPEbarrel with white opaque LDPE plunger and white opaque LDPE cap. Cleaning towels (smooth, white crepe paper impregnated with isopropylalcohol/benzalkonium chloride) individually wrapped. Cardboard boxes of 12 syringes and 12 cleaning towels.

Prefilled 8 g single-dose intramammary syringe consisting of white opaque LDPEbarrel with white opaque LDPE plunger and white opaque LDPE cap.Cleaning towels (smooth, white crepe paper impregnated with isopropylalcohol/benzalkonium

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Industrial Veterinaria S.A.

Marketing authorisation date:

24/05/2018

Manufacturing sites for batch release:

aniMedica GmbH

Industrial Veterinaria S.A.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

3663 ESP

Date of authorisation status change:

26/05/2018

Reference member state:

Ireland

Procedure number:

IE/V/0480/001

Concerned member states:

Austria Belgium Croatia Cyprus France Germany Greece Hungary Italy Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland) 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

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

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