Cobactan LC, 75 mg, intramammary ointment for lactating cattle

Authorised

- Cefquinome sulfate

Product identification

Medicine name:
Cobactan LC, 75 mg, intramammary ointment for lactating cattle

Active substance:
Cefquinome sulfate

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Cefquinome sulfate
88.92 milligram(s) / 8.00 gram(s)

Pharmaceutical form:

Intramammary ointment

Withdrawal period by route of administration:

Intramammary use:

- **Cattle**
  - Meat and offal. 4 day
  - Milk. 5 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51DE90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Package description:

White opaque polyethylene syringes and cleaning towels in paper aluminium copolymer laminate sachet.Packs of 3 syringes and cleaning towels.
White opaque polyethylene syringes and cleaning towels in paper aluminium copolymer laminate sachet.Packs of 15 syringes and cleaning towels.
White opaque polyethylene syringes and cleaning towels in paper aluminium copolymer laminate sachet.Packs of 20 syringes and cleaning towels.
White opaque polyethylene syringes and cleaning towels in paper aluminium copolymer laminate sachet.Packs of 24 syringes and cleaning towels.

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:
Intervet Nederland B.V.

Marketing authorisation date:
31/07/1998

Manufacturing sites for batch release:
Intervet International GmbH
Intervet International B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 9468

Date of authorisation status change:
27/01/2022

Reference member state:
Ireland

Procedure number:
IE/V/0467/001

Concerned member states:
Austria Belgium Denmark France Germany Greece Hungary Italy Luxembourg Netherlands Poland Portugal Romania Slovakia Spain United Kingdom (Northern Ireland)

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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Summary of Product Characteristics