

# VIRBACTAN 150 MG INTRAMAMMARY OINTMENT

Authorised

- Cefquinome sulfate

## Product identification

**Medicine name:**

VIRBACTAN 150 MG INTRAMAMMARY OINTMENT

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**Active substance:**

Cefquinome sulfate

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**Target species:**

Cattle (dry cow)

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**Route of administration:**

Intramammary use

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## Product details

**Active substance and strength:**

Cefquinome sulfate

177.80 milligram(s) / 1.00 Syringe

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**Pharmaceutical form:**

Intramammary ointment

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**Withdrawal period by route of administration:****Intramammary use:**

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**Cattle (dry cow)**

- Meat and offal. 2 day

- Milk. 36 day

Milk: 36 days after treatment when dry period is 5 weeks or less.

- Milk. 1 day

Milk: 1 day after calving when dry period is more than 5 weeks

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ51DE90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Slovenia

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**Available in:**

Slovenia

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**Package description:**

Box of 1 sachet of 4 applicators and 4 cleaning towels

Box of 30 sachets of 4 applicators and 120 cleaning towels

Box of 15 sachets of 4 applicators and 60 cleaning towels

Box of 6 sachets of 4 applicators and 24 cleaning towels

Box of 5 sachets of 4 applicators and 20 cleaning towels

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Virbac

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**Marketing authorisation date:**

17/01/2006

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**Manufacturing sites for batch release:**

Haupt Pharma Latina S.r.l.

Virbac

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**Responsible authority:**

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

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**Authorisation number:**

MR/V/0358/001

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**Date of authorisation status change:**

17/01/2006

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**Reference member state:**

France

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**Procedure number:**

FR/V/0148/001

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**Concerned member states:**

Austria Belgium Cyprus Czechia Estonia Germany Greece Ireland Italy  
Latvia Lithuania Luxembourg Netherlands Poland Portugal Slovakia Slovenia  
Spain United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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.

### Combined File of all Documents