

VIRBACTAN 150 MG INTRAMAMMARY OINTMENT

Authorised

- Cefquinome sulfate

Product identification

Medicine name:

VIRBACTAN 150 MG INTRAMAMMARY OINTMENT

Active substance:

Cefquinome sulfate

Target species:

Cattle (dry cow)

Route of administration:

Intramammary use

Product details

Active substance and strength:

Cefquinome sulfate

177.80 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary ointment

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51DE90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Available in:

Slovakia

Package description:

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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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:

Virbac

Marketing authorisation date:

1/04/2005

Manufacturing sites for batch release:

Haupt Pharma Latina S.r.l.

Virbac

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/009/MR/05-S

Date of authorisation status change:

1/04/2005

Reference member state:

France

Procedure number:

FR/V/0148/001

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)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

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