

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:

Cyprus

Available in:

Cyprus

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

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:

30/05/2005

Manufacturing sites for batch release:

Haupt Pharma Latina S.r.l.

Virbac

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

CY00089V

Date of authorisation status change:

4/02/2015

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

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