

Fatroximin DC 100 mg - Salbe zur intramammären Anwendung für Rinder

Authorised

- Rifaximin

Product identification

Medicine name:

Fatroximin DC 100 mg - Salbe zur intramammären Anwendung für Rinder

Active substance:

Rifaximin

Target species:

Cattle (dry cow)

Route of administration:

Intramammary use

Product details

Active substance and strength:

Rifaximin

100.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary ointment

Withdrawal period by route of administration:

Intramammary use:

-

Cattle (dry cow)

- Meat and offal. 0 day

1 Euterinjektor (100 mg Rifaximin) pro Euterviertel, einmalig in alle 4 Euterviertel

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51XX01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria

Package description:

Available only in [German](#)

Available only in [German](#)

Available only in [German](#)

Available only in [German](#)

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:

Fatro S.p.A.

Marketing authorisation date:

29/12/1998

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

8-00374

Date of authorisation status change:

29/12/1998

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

Documents

Package Leaflet

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Summary of Product Characteristics

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.