

# FATROXIMIN D.C. 100 mg/5 ml intramamálna masť

Authorised

- Rifaximin

## Product identification

**Medicine name:**

FATROXIMIN D.C. 100 mg/5 ml intramamálna masť

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**Active substance:**

Rifaximin

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**Target species:**

Cattle (dry cow)

Buffalo (female)

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**Route of administration:**

Intramammary use

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## Product details

**Active substance and strength:**

Rifaximin

100.00 milligram(s) / 1.00 Applicator

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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. 0 day

Meat and offal: without withdrawal period, The udder must be excluded from human consumption.

- Milk. 0 day

Milk: 0 days after calving if the length of dry standing is equal to or longer than 35 days. 35 days after administration if the duration of dry standing is less than 35 days.

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**Buffalo (female)**

- Meat and offal. 0 day

Meat and offal: without withdrawal period, The udder must be excluded from human consumption.

- Milk. 0 day

Milk: 0 days after calving if the length of dry standing is equal to or longer than 35 days. 35 days after administration if the duration of dry standing is less than 35 days.

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

QJ51XX01

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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:**

Slovakia

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**Package description:**

Available only in [Slovak](#)

Available only in [Slovak](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Fatro S.p.A.

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**Marketing authorisation date:**

23/12/1997

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**Manufacturing sites for batch release:**

Fatro S.p.A.

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/0638/97-S

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**Date of authorisation status change:**

23/12/1997

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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

Combined File of all Documents

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