

FATROXIMIN 7,5 mg/ g intrauterinná a intravaginálna pena

Authorised

- Rifaximin

Product identification

Medicine name:

FATROXIMIN 7,5 mg/ g intrauterinná a intravaginálna pena

Active substance:

Rifaximin

Target species:

Cattle (cow)

Horse (mare)

Route of administration:

Intrauterine use

Vaginal use

Product details

Active substance and strength:

Rifaximin

100.50 milligram(s) / 1.00 Container

Pharmaceutical form:

Intrauterine foam

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

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Cattle (cow)

- Meat and offal. 0 day Meat and offal: zero days.

- Milk. 0 day Milk: 0 hours.

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Horse (mare)

- Meat and offal. 0 day

Meat and offal: zero days. Do not use in mares whose meat is intended for human consumption.

- Milk. 0 day Milk: zero hours

Vaginal use:

-

Cattle (cow)

- Meat and offal. 0 day Meat and offal: zero days.

- Milk. 0 day Milk: zero hours.

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Horse (mare)

- Meat and offal. 0 day

Meat and offal: zero days. Do not use in mares whose meat is intended for human consumption.

- Milk. 0 day Milk: zero hours.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG51AA06

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 Slovak

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Fatro S.p.A.

Marketing authorisation date:

23/12/1997

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/0636/97-S

Date of authorisation status change:

23/12/1997

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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