Fatroximin Intrauterine Foam 0,75 g/100 g Aerozol

Authorised

- Rifaximin

Product identification

Medicine name:
Fatroximin Intrauterine Foam 0,75 g/100 g Aerozol

Active substance:
Rifaximin

Target species:
Cattle
Horse

Route of administration:
Vaginal use
Intrauterine use

Product details

Active substance and strength:
Rifaximin
0.75 gram(s) / 100.00 gram(s)

Pharmaceutical form:

This information is not available for this product.

Withdrawal period by route of administration:

Vaginal use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Horse**
  - All relevant tissues. no withdrawal period

Not authorised for use in horses intended for human consumption.

Intrauterine use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Horse**
  - All relevant tissues. no withdrawal period

Not authorised for use in horses intended for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG51AA06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Available in:
Poland

Package description:
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
20/06/2000

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1020

Date of authorisation status change:
20/06/2000

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

Documents

Labelling

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Package Leaflet

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