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

This document does not exist in this language (English). You can find it in another language below.