Fatroximin, 300mg, Intrauterinní tableta

Authorised

- Rifaximin

Product identification

Medicine name:
Fatroximin, 300mg, Intrauterinní tableta

Active substance:
Rifaximin

Target species:
Horse (mare)
Buffalo (female)
Cattle (cow)

Route of administration:
Vaginal use
Intrauterine use

Product details

Active substance and strength:
Rifaximin
300.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Intrauterine tablet

Withdrawal period by route of administration:
Vaginal use:

- **Horse (mare)**
  - Meat and offal. no withdrawal period

Nepoužívat u koní, jejichž maso a mléko je určeno pro lidskou spotřebu.,

  - Milk. no withdrawal period

Nepoužívat u koní, jejichž maso a mléko je určeno pro lidskou spotřebu.,

- **Buffalo (female)**
  - Meat and offal. 0 day
  - Milk. 0 hour

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 hour

Intrauterine use:

- **Horse (mare)**
  - Meat and offal. no withdrawal period

Nepoužívat u koní, jejichž maso a mléko je určeno pro lidskou spotřebu.,

  - Milk. no withdrawal period

Nepoužívat u koní, jejichž maso a mléko je určeno pro lidskou spotřebu.,

- **Buffalo (female)**
  - Meat and offal. 0 day

- Milk. 0 hour

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG51AA06

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech
Available only in Czech

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:
19/09/1997

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/841/97-C

Date of authorisation status change:
23/05/2013

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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.