FATROXIMIN 300 mg intrauterinné tablety

Authorised

- Rifaximin

Product identification

Medicine name:
FATROXIMIN 300 mg intrauterinné tablety

Active substance:
Rifaximin

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

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:

- **Cattle (cow)**
  - Milk. 0 day
    Milk: zero hours.
  - Meat and offal. 0 day
    Meat and offal: zero days.
- **Buffalo (female)**
  - Milk. 0 day
    Milk: zero hours.
  - Meat and offal. 0 day
    Meat and offal: zero days.
- **Horse (mare)**
  - Milk. 0 day
    Milk: zero hours.
  - Meat and offal. 0 day

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

Intrauterine use:

- **Cattle (cow)**
  - Meat and offal. 0 day
    Meat and offal: zero days.
  - Milk. 0 day
    Milk: zero hours.
- **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: 0 hours.

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

Meat and offal: zero days.

  - Milk. 0 day

Milk: zero hours.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG51AA06

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

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Authorisation status:
Valid

Authorised in:
Slovakia

Available in:
Slovakia

Package description:
Available only in Slovak
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:
96/0637/97-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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