Fatroximin, 100 mg/13,4g, intrauteriinvaht hobustele, kelle liha ei tarvitata inimtoiduks ja veistele

Authorised

- Rifaximin

Product identification

Medicine name:
Fatroximin, 100 mg/13,4g, intrauteriinvaht hobustele, kelle liha ei tarvitata inimtoiduks ja veistele

Active substance:
Rifaximin

Target species:
Horse (non food-producing)
Cattle

Route of administration:
Intrauterine use

Product details

Active substance and strength:

Rifaximin
100.00 milligram(s) / 13.40 gram(s)

Pharmaceutical form:
Intrauterine foam

Withdrawal period by route of administration:
Intrauterine use:

- **Horse (non food-producing)**
  - Milk. no withdrawal period

Mitte manustada märadele, kelle liha, söödavaid kudesid ja piima tarvitatakse inimtoiduks.

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour Piim: 0 tundi (0 lüpsi).

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG51AA06

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

Authorisation status:
Valid

Authorised in:
Estonia

Package description:
Available only in Estonian

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:
3/10/2002

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

Responsible authority:
State Agency Of Medicines

Authorisation number:
1107

Date of authorisation status change:
3/10/2002

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

Documents

Summary of Product Characteristics

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