Fatroximin, 2 mg/ml nahasprei, lahus veistele, hobustele, lammastele, kitsedele, sigadele, küülikutele, koertele, kassidele.

Authorised

- Rifaximin

Product identification

Medicine name:
Fatroximin, 2 mg/ml nahasprei, lahus veistele, hobustele, lammastele, kitsedele, sigadele, küülikutele, koertele, kassidele.

Active substance:
Rifaximin

Target species:
Cattle
Horse
Pig
Sheep
Goat
Rabbit
Dog
Cat

Route of administration:

Cutaneous use

Product details

Active substance and strength:
Rifaximin
2.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Cutaneous spray, solution

Withdrawal period by route of administration:
Cutaneous use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
- **Goat**
  - Meat and offal. 0 day
- **Rabbit**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD06AX11

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:
1082

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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