

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
- Milk. 0 day

-

Pig

- Meat and offal. 0 day

-

Sheep

- Meat and offal. 0 day
- Milk. 0 day

-

Goat

- Meat and offal. 0 day
- Milk. 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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