Fatroximin Dry Cow, 100 mg/5ml intramammaarsalv kinnislehmadele

Authorised

- Rifaximin

Product identification

Medicine name:
Fatroximin Dry Cow, 100 mg/5ml intramammaarsalv kinnislehmadele

Active substance:
Rifaximin

Target species:
Cattle (dry cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Rifaximin
100.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary ointment

Withdrawal period by route of administration:

Intramammary use:

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

Mitte tarvitada ravitud loomade udarat inimtoiduks.

0 päeva, kui kinnisperiood on pikem kui 42 päeva; 15 lüpsikorda, kui kinnisperiood on lühem kui 42 päeva.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51XX01

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:
1110

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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