FATROXIMIN, ενδομαστική αλοιφή

Authorised

- Rifaximin

Product identification

Medicine name:
FATROXIMIN, ενδομαστική αλοιφή

Active substance:
Rifaximin

Target species:
Cattle (cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Rifaximin
0.10 gram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary ointment

Withdrawal period by route of administration:
Intramammary use:

-

Cattle (cow)

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51XX01

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

Authorisation status:
Valid

Authorised in:
Greece

Available in:
Greece

Package description:
Available only in Greek
Available only in Greek
Available only in Greek

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
24/02/1991

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

Responsible authority:
National Organization For Medicines

Authorisation number:
65434/14-09-2012/K-0036502

Date of authorisation status change:
17/06/2024

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.
