FATROXIMIN DRY COW, intramaminė suspensija

Authorised

- Rifaximin

Product identification

Medicine name:
FATROXIMIN DRY COW, intramaminė suspensija

Active substance:
Rifaximin

Target species:
Cattle (cow)

Route of administration:
Intramammary use

Product details

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

Pharmaceutical form:
Intramammary suspension

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51XX01

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

Authorisation status:
Valid

Authorised in:
Lithuania

Available in:
Lithuania

Package description:
Available only in Lithuanian
Available only in Lithuanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
8/07/2006

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

Responsible authority:
State Food And Veterinary Service

Authorisation number:

LT/2/01/1284/001-002

Date of authorisation status change:
8/07/2006

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

Documents

Combined File of all Documents

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