Fatroximin 20 mg/ml pomada instramamária para bovinos

Authorised

- Rifaximin

Product identification

Medicine name:
Fatroximin 20 mg/ml pomada instramamária para bovinos

Active substance:
Rifaximin

Target species:
Cattle (dairy cow at drying-off)

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 (dairy cow at drying-off)**
  - Meat and offal. 0 day

Não utilizar a carne do ubere para consumo humano. Leite: Zero dias após o parto quando o período de secagem é igual ou superior a 35 dias, 35 dias após o tratamento quando o período de secagem é inferior a 35 dias

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Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51XX01

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

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Authorisation status:
Valid

Authorised in:
Portugal

Available in:
Portugal

Package description:
Available only in Portuguese
Available only in Portuguese

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:
4/12/1992

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

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
50870

Date of authorisation status change:
27/02/2014

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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