Fatroximin D.C. 100 mg/5 ml zawiesina dowymieniowa dla bydła

Authorised

- Rifaximin

Product identification

Medicine name:
Fatroximin D.C. 100 mg/5 ml zawiesina dowymieniowa dla bydła

Active substance:
Rifaximin

Target species:
Cattle

Route of administration:
Intramammary use

Product details

Active substance and strength:
Rifaximin
100.00 milligram(s) / 5.00 millilitre(s)

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle**
  - Milk. 2 day
  - Milk. 35 day
  - 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:
Poland

Available in:
Poland

Package description:
Available only in Polish
Available only in Polish

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:
20/06/2000

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

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1021

Date of authorisation status change:
20/06/2000

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

Documents

Package Leaflet

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Labelling

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

Summary of Product Characteristics

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