FATROXIMIN DC 100 mg/5 ml intramammális kenőcs szarvasmarhák és bivalyok részére A.U.V.

Authorised

- Rifaximin

Product identification

Medicine name:
FATROXIMIN DC 100 mg/5 ml intramammális kenőcs szarvasmarhák és bivalyok részére A.U.V.

Active substance:
Rifaximin

Target species:
Cattle
Buffalo

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**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Buffalo**
  - 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:
Hungary

Package description:
Available only in Hungarian
Available only in Hungarian
Available only in Hungarian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
20/11/2019

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

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
20/11/2019

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

Documents

Summary of Product Characteristics

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Package Leaflet

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