Fatroximin 100 mg/13,4 g putas intrauterīnai lietošanai govīm un ķēvēm

Authorised

- Rifaximin

Product identification

Medicine name:
Fatroximin 100 mg/13,4 g putas intrauterīnai lietošanai govīm un ķēvēm

Active substance:
Rifaximin

Target species:
Horse (mare)
Cattle (cow)

Route of administration:
Intrauterine use

Product details

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

Pharmaceutical form:
Intrauterine foam

Withdrawal period by route of administration:

Intrauterine use:

- **Horse (mare)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Cattle (cow)**
  - Milk. 0 day
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG51AA06

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

Authorisation status:
Valid

Authorised in:
Latvia

Package description:
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
5/07/2001

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

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/01/1371

Date of authorisation status change:
5/07/2001

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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