FATROXIMIN 7,5 mg/ml emulsão intra-uterina para bovinos e equinos

Authorised

- Rifaximin

Product identification

Medicine name:
FATROXIMIN 7,5 mg/ml emulsão intra-uterina para bovinos e equinos

Active substance:
Rifaximin

Target species:
Cattle
Horse

Route of administration:
Intrauterine use

Product details

Active substance and strength:
Rifaximin
7.50 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Intrauterine foam

Withdrawal period by route of administration:
Intrauterine use:

- **Cattle**
  - Meat and offal. 0 day — Carne e vísceras: zero dias Leite: zero dias
  - Milk. 0 day — Carne e vísceras: zero dias Leite: zero dias
- **Horse**
  - Meat and offal. 0 day — Carne e vísceras: zero dias Leite: zero dias

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG51AA06

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

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:
50871

Date of authorisation status change:
1/03/2014

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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