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FATROXIMIN TOPIC, 2.94mg/g, Kožní sprej, roztok

Authorised

- Rifaximin

Product identification

Medicine name:
FATROXIMIN TOPIC, 2.94mg/g, Kožní sprej, roztok

Active substance:
Rifaximin

Target species:
Cat
Horse
Goat
Rabbit
Sheep
Pig
Cattle
Dog

Route of administration:
Cutaneous use

Product details

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

Pharmaceutical form:
Cutaneous spray, solution

Withdrawal period by route of administration:
Cutaneous use:

- **Horse**
  - Meat and offal. 0 day
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Rabbit**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Cattle**
  - Meat and offal. 0 day

- Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD06AX11

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

Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech

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:
8/12/1998

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

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/148/98-C

Date of authorisation status change:
23/05/2013

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.

Labelling

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