FATROXIMIN TOPIC 2,94 mg/g kožný sprej, roztok

Authorised

- Rifaximin

Product identification

Medicine name:
FATROXIMIN TOPIC 2,94 mg/g kožný sprej, roztok

Active substance:
Rifaximin

Target species:
Cattle
Sheep
Goat
Pig
Horse
Rabbit
Dog
Cat

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:

- **Cattle**
  - All relevant tissues. 0 day — Without withdrawal period
- **Sheep**
  - All relevant tissues. 0 day — Without withdrawal period
- **Goat**
  - All relevant tissues. 0 day — Without withdrawal period
- **Pig**
  - All relevant tissues. 0 day — Without withdrawal period
- **Horse**
  - All relevant tissues. 0 day — Without withdrawal period
- **Rabbit**
  - All relevant tissues. 0 day — Without withdrawal period
- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD06AX11

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

Authorisation status:
Valid

Authorised in:
Slovakia

Available in:
Slovakia

Package description:
Available only in Slovak

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
6/09/2001

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

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

Authorisation number:

96/074/01-S

Date of authorisation status change:
6/09/2001

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

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