Rivanol unguent

Authorised

- Ethacridine lactate

Product identification

Medicine name:
Риванол унгвент
Rivanol unguent

Active substance:
Ethacridine lactate

Target species:
Horse
Dog

Route of administration:
External use

Product details

Active substance and strength:
Ethacridine lactate
1.00 gram(s) / 100.00 gram(s)

Pharmaceutical form:
Ointment

Withdrawal period by route of administration:

External use:

- **Horse**
  - Meat and offal. no withdrawal period
- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD08AA01

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian

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:
Farma Vet OOD

Marketing authorisation date:
12/07/2012

Manufacturing sites for batch release:
Farma Vet OOD

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1811

Date of authorisation status change:
12/07/2012

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 and Labelling

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