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

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

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

Package Leaflet and Labelling

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