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Equibactin vet. 333 mg/g + 67 mg/g oral paste for horses

Authorised

- Trimethoprim
- Sulfadiazine

Product identification

Medicine name:
Equibactin vet. 333 mg/g + 67 mg/g oral paste for horses

Active substance:
Trimethoprim

Sulfadiazine

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Trimethoprim
66.70 milligram(s) / 1.00 gram(s)

Sulfadiazine

333.30 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral paste

Withdrawal period by route of administration:
Oral use:

- **Horse**
  - Meat and offal. 14 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01EW10

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

Authorisation status:
Valid

Authorised in:
Denmark

Package description:
5 pre-filled multi-dose (Low Density) polyethylene syringes containing 45 g paste with adjustable screw ring closed with a (Low Density) polyethylene cap, packed in a cardboard box.
1 pre-filled multi-dose (Low Density) polyethylene syringe containing 45 g paste with adjustable screw ring closed with a (Low Density) polyethylene cap, packed in a cardboard box.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Le Vet. B.V.

Marketing authorisation date:
9/06/2008

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Danish Medicines Agency

Authorisation number:
39786

Date of authorisation status change:
9/06/2008

Reference member state:
Netherlands

Procedure number:
NL/V/0123/001

Concerned member states:
Austria Belgium Denmark Finland France Germany Greece Hungary Iceland Ireland Italy Luxembourg Norway Portugal Spain Sweden

United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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Combined File of all Documents