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Flukiver Combi 50 mg/ml + 75 mg/ml oral suspension

Authorised

- Mebendazole
- Closantel sodium dihydrate

Product identification

Medicine name:
Flukiver Combi 50 mg/ml + 75 mg/ml oral suspension

Active substance:
Mebendazole

Closantel sodium dihydrate

Target species:
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Mebendazole
75.00 milligram(s) / 1.00 millilitre(s)

Closantel sodium dihydrate

54.37 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:
Oral use:

- **Sheep**
  - Meat and offal. 65 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52A

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

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
Container: High density polyethylene bottles of 1 litre Closure: High density polyethylene screw cap
Container: High density polyethylene bottles of 2.5 litresClosure: High density polyethylene screw cap
Container: High density polyethylene bottles of 5 litresClosure: High density polyethylene screw cap

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:
Elanco GmbH

Marketing authorisation date:
30/06/2009

Manufacturing sites for batch release:
Lusomedicamenta Sociedade Tecnica Farmaceutica S.A.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401206.00.00

Date of authorisation status change:
22/12/2014

Reference member state:
Ireland

Procedure number:
IE/V/0222/001

Concerned member states:
Austria Belgium France Germany Iceland Italy Netherlands

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

Documents

Combined File of all Documents

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