ALBENDAZOL suspension

Authorised

- Albendazole

Product identification

Medicine name:
ALBENDAZOL suspension

Active substance:
Albendazole

Target species:
Cattle
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Albendazole
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 27 day
  - Milk. 72 hour
- **Sheep**
  - Meat and offal. 8 day

Not authorised for use in animals producing milk for human consumption

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AC11

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Farma Vet OOD

Marketing authorisation date:
24/05/2017

Manufacturing sites for batch release:
Farma Vet OOD

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2736

Date of authorisation status change:
24/05/2017

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

Documents

Summary of Product Characteristics

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Labelling

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

Package Leaflet

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