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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 Ltd.

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

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