ALBENDAZOL 300 mg, 600 mg and 1200 mg

Authorised

- Albendazole
- Albendazole
- Albendazole

Product identification

Medicine name:
ALBENDAZOL 300 mg, 600 mg and 1200 mg

Active substance:
Albendazole
Albendazole
Albendazole

Target species:
Cattle
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:

Albendazole
1200.00 milligram(s) / 1.00 Tablet

Albendazole
600.00 milligram(s) / 1.00 Tablet

Albendazole
300.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 14 day
  - Milk. 72 hour

мляко - 72 часа след последното третиране

- **Sheep**
  - Meat and offal. 7 day

мляко - не се разрешава за употреба при овце, чието мляко е предназначено за консумация от хора

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AC11

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

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Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian

Available only in Bulgarian
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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 Sis OOD

Marketing authorisation date:
28/10/2007

Manufacturing sites for batch release:
Golash Pharma OOD

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1971

Date of authorisation status change:
6/03/2013

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.