Bovitak 2,265%, suspensie orală pentru bovine și ovine

Not authorised

- Oxfendazole

Product identification

Medicine name:
Bovitak 2,265%, suspensie orală pentru bovine și ovine

Active substance:
Oxfendazole

Target species:
Cattle
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Oxfendazole
2.27 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Cattle**

- Meat and offal. 14 day

Animals intended for human consumption should not be slaughtered during treatment.

- Milk. 84 hour

Animals intended for human consumption should not be slaughtered during treatment.

- **Sheep**

- Meat and offal. 21 day

Animals intended for human consumption should not be slaughtered during treatment.

- Milk. no withdrawal period

Use during lactation in sheep producing milk for human consumption is not permitted.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AC02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Surrendered

Authorised in:

Romania

Package description:

Available only in Romanian

Available only in Romanian

Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Richter Pharma S.R.L.

Marketing authorisation date:
26/07/2015

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
200207

Date of authorisation status change:
6/03/2025

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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