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BULMECTIN 2 mg/g, premix pentru furaj medicamentat pentru bovine, oi și cai

Authorised

- Abamectin

Product identification

Medicine name:
BULMECTIN 2 mg/g, premix pentru furaj medicamentat pentru bovine, oi și cai

Active substance:
Abamectin

Target species:
Cattle
Sheep
Horse (non-food-producing)

Route of administration:
Oral use

Product details

Active substance and strength:
Abamectin
2.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Premix for medicated feeding stuff

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 21 day

Nu se administrează la animalele a căror lapte este destinat consumului uman.

- **Sheep**
  - Meat and offal. 21 day

Nu se administrează la animalele a căror lapte este destinat consumului uman.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA02

Legal status of supply:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Romania

Package description:
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian
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:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Biovet AD

Marketing authorisation date:
27/06/2001

Manufacturing sites for batch release:
Biovet AD

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

Authorisation number:
140010

Date of authorisation status change:
2/09/2025

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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