PANAFEN

Authorised

- Fenbendazole

Product identification

Medicine name:
PANAFEN

Active substance:
Fenbendazole

Target species:
Cattle
Horse
Goat
Sheep

Route of administration:
Buccal use

Product details

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

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Buccal use:

- **Cattle**
  - Meat and offal. 9 day
  - Milk. 5 day
- **Horse**
  - Meat and offal. no withdrawal period

It is not administered to horses whose meat is intended for human consumption.

- **Goat**
  - Meat and offal. 18 day
  - Milk. 9 day
- **Sheep**
  - Meat and offal. 18 day
  - Milk. 9 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AC13

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Romania

Available 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

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Pasteur Filiala Filipesti S.A.

Marketing authorisation date:
2/07/2013

Manufacturing sites for batch release:
Pasteur Filiala Filipesti S.A.

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

Authorisation number:
190152

Date of authorisation status change:
26/08/2025

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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