Panacur AquaSol 200 mg/ml - Oral suspension

Authorised

- Fenbendazole

Product identification

Medicine name:
Panacur AquaSol 200 mg/ml - Oral suspension

Active substance:
Fenbendazole

Target species:
Chicken
Pig

Route of administration:
In drinking water use

Product details

Active substance and strength:
Fenbendazole
200.00 milligram(s) / 1.00 Bottle

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

In drinking water use:

- **Chicken**
  - Meat and offal. 6 day

6 days for 1 mg fenbendazole/kg dose; 9 days for 2 mg fenbendazole/kg dose

  - Eggs. 0 day Zero days

- **Pig**
  - Meat and offal. 4 day 4 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AC13

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Belgium , Croatia , Denmark , Finland , France , Germany , Hungary , Ireland , Italy , Latvia , Lithuania , Netherlands , Norway , Poland , Slovenia , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:4 L

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:1 L

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:
Intervet International B.V.

Marketing authorisation date:
9/12/2011

Manufacturing sites for batch release:
Intervet Productions S.A.

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
9/12/2011

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

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