Pulmotil AC 250 mg/mL concentrate for oral solution for use in drinking water or milk replacer for chickens, turkeys, pigs and calves

Authorised

- Tilmicosin

Product identification

Medicine name:
Pulmotil AC 250 mg/mL concentrate for oral solution for use in drinking water or milk replacer for chickens, turkeys, pigs and calves
Pulmotil AC 250 mg/ml concentraat voor orale oplossing voor gebruik in drinkwater voor kippen, varkens, kalkoenen, of in kunstmelk voor kalveren

Active substance:
Tilmicosin

Target species:
Chicken
Turkey
Pig
Cattle (pre-ruminant)

Route of administration:

Oral use

Product details

Active substance and strength:
Tilmicosin
250.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Concentrate for oral solution

Withdrawal period by route of administration:
Oral use:

- **Chicken**
  - Meat and offal. 12 day

Not authorised for use in laying birds producing eggs for human consumption. Do not use within 14 days of the start of the laying period

- **Turkey**
  - Meat and offal. 19 day

Not authorised for use in laying birds producing eggs for human consumption. Do not use within 14 days of the start of the laying period

- **Pig**
  - Meat and offal. 14 day

- **Cattle (pre-ruminant)**
  - Meat and offal. 42 day

Not authorised for use in animals producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA91

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

Authorisation status:
Valid

Authorised in:
Netherlands

Available in:
Netherlands

Package description:
Polyethylene naphthalate amber coloured bottle containing 960 mL of veterinary medicinal product, with a polypropylene screw top and polyethylene/aluminium/polyethylene terephthalate seal.
Polyethylene naphthalate amber coloured bottle containing 240 mL of veterinary medicinal product, with a polypropylene screw top and polyethylene/aluminium/polyethylene terephthalate seal.

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:
Elanco GmbH

Marketing authorisation date:
19/04/2002

Manufacturing sites for batch release:
Elanco France S.A.S.

Responsible authority:

Medicines Evaluation Board

Authorisation number:
REG NL 9972

Date of authorisation status change:
24/02/2022

Reference member state:
Italy

Procedure number:
IT/V/0102/001

Concerned member states:
Belgium Greece Luxembourg Netherlands Portugal Spain

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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