

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

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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

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Pig

- Meat and offal. 14 day

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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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