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 πυκνό διάλυμα για την παρασκευή πόσιμου διαλύματος προς χρήση σε πόσιμο νερό ή υποκατάστατο γάλακτος για όρνιθες, ινδόρνιθες, χοίρους και μόσχους

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:
Greece

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:
17/05/2010

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

Responsible authority:
National Organization For Medicines

Authorisation number:
33007/18-05-2010/K-0119902

Date of authorisation status change:
17/05/2010

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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.