

TILMOVET 25% oral solution for chickens, turkeys, pigs and calves

Authorised

- Tilmicosin

Product identification

Medicine name:

TILMOVET 25% oral solution for chickens, turkeys, pigs and calves

Active substance:

Tilmicosin

Target species:

Chicken (broiler)

Chicken (chick, for replacement)

Chicken (chick)

Turkey

Pig

Cattle (calf)

Route of administration:

In drinking water use

Product details

Active substance and strength:

Tilmicosin

250.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral solution

Withdrawal period by route of administration:

In drinking water use:

•

Chicken (broiler)

- Meat and offal. 12 day

Яйца: Не се разрешава употребата при птици-носачки, чиито яйца са предназначени за човешка консумация.

•

Chicken (chick, for replacement)

- Meat and offal. 12 day

•

Chicken (chick)

- Meat and offal. 12 day

•

Turkey

- Meat and offal. 19 day

Яйца: Не се разрешава употребата при птици, които произвеждат или са предназначени да произвеждат яйца за консумация от хора. Да не се прилага в рамките на 14 дни от началото на яйценосния период

•

Pig

- Meat and offal. 14 day

•

Cattle (calf)

- Meat and offal. 42 day

Не се разрешава за употреба при животни, чието мляко е предназначено за консумация от хора

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Package description:

Available only in [Bulgarian](#)

Available only in [Bulgarian](#)

Available only in [Bulgarian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Biovet AD

Marketing authorisation date:

21/06/2006

Manufacturing sites for batch release:

Biovet AD

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-1549

Date of authorisation status change:

8/06/2023

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

Documents

Summary of Product Characteristics

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

Package Leaflet and Labelling

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