

TYLOGRAN, 1000 mg/g, granules for use in drinking water/milk for cattle (calves), pigs, chickens and turkeys

Authorised

- Tylosin tartrate

Product identification

Medicine name:

TYLOGRAN, 1000 mg/g, granules for use in drinking water/milk for cattle (calves), pigs, chickens and turkeys

Active substance:

Tylosin tartrate

Target species:

Pig

Chicken

Cattle (calf)

Turkey

Route of administration:

In drinking water use

Oral use

Product details

Active substance and strength:

Tylosin tartrate

1.10 gram(s) / 1.10 gram(s)

Pharmaceutical form:

Granules for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

-

Pig

- Meat and offal. 1 day

-

Chicken

- Meat and offal. 1 day

- Egg. 0 day

-

Cattle (calf)

- Meat and offal. 12 day

-

Turkey

- Meat and offal. 2 day

- Egg. 0 day

Oral use:

-

Pig

- Meat and offal. 1 day

-

Chicken

- Meat and offal. 1 day

- Egg. 0 day

•

Cattle (calf)

- Meat and offal. 12 day

•

Turkey

- Meat and offal. 2 day

- Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Available in:

Poland

Package description:

Securitainer: white polypropylene cylindrical container provided with a low density polyethylene lid. The securitainer contains 550 g of product.

Composite can: hardboard can provided with an inner lining of aluminium paper (polyethylene polyethylene terephthalate coated) and a seamed tin plate bottom, closed with a low density polyethylene lid. The can contains 550 g of product.

Securitainer: white polypropylene cylindrical container provided with a low density polyethylene lid. The securitainer contains 100 g of product.

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Securitainer: white polypropylene cylindrical container provided with a low density polyethylene lid. The securitainer contains 800 g of product.

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Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Dopharma Research B.V.

Marketing authorisation date:

3/11/2015

Manufacturing sites for batch release:

Dopharma B.V.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

2480

Date of authorisation status change:

3/11/2015

Reference member state:

Netherlands

Procedure number:

NL/V/0189/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland
France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania
Luxembourg Norway Poland Portugal Romania Slovakia Sweden

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

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