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TYLUCYL 200 MG/ML SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Authorised

- Tylosin

Product identification

Medicine name:
TYLUCYL 200 MG/ML SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Active substance:
Tylosin

Target species:
Cattle
Pig

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Tylosin
200000.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 108 hour
- **Pig**
  - Meat and offal. 16 day

Intravenous use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 108 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01FA90

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

Authorisation status:
Valid

Authorised in:
Greece

Package description:
50 ml colourless type II glass vials closed with a bromobutyl rubber stopper and aluminium cap. One vial per carton.
250 ml colourless type II glass vials closed with a bromobutyl rubber stopper and aluminium cap. One vial per carton.

100 ml colourless type II glass vials closed with a bromobutyl rubber stopper and aluminium cap. One vial per carton.

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:
Vetoquinol S.A.

Marketing authorisation date:
15/09/2016

Manufacturing sites for batch release:
Vetoquinol S.A.

Responsible authority:
National Organization For Medicines

Authorisation number:
74770/03-08-2021/K-0213601

Date of authorisation status change:
2/08/2021

Reference member state:
France

Procedure number:
FR/V/0289/001

Concerned member states:
Austria Belgium Croatia Cyprus Czechia Denmark Estonia Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Malta Netherlands Poland Portugal Romania Slovakia Slovenia Spain Sweden

United Kingdom (Northern Ireland)

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.

Combined File of all Documents

eu-puar-frv0289001-mr-rpe214-en.pdf