Toltrazuril

Authorised

- Toltrazuril

Product identification

Medicine name:
Toltrazuril
Coccinox 25 mg/ml Oplossing voor gebruik in drinkwater
Coccinox 25 mg/ml Solution pour administration dans l'eau de boisson
Coccinox 25 mg/ml Lösung zum Eingeben über das Trinkwasser

Active substance:
Toltrazuril

Target species:
Turkey
Chicken

Route of administration:
Oral use

Product details

Active substance and strength:
Toltrazuril
25.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for use in drinking water

Withdrawal period by route of administration:

Oral use:

- **Turkey**
  - Meat and offal. 16 day

- **Chicken**
  - Meat and offal. 16 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP51AJ01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Package description:

1 x 50 ml - Type III amber glass bottle of 50 ml closed with tamper-evident HDPE screw caps with ring and colourless LDPE syringe insert. Each bottle is individually packed in a cardboard box

1 x 10 ml - Type III amber glass bottle of 10 ml closed with tamper-evident HDPE screw caps with ring and colourless LDPE syringe insert. Each bottle is individually packed in a cardboard box

10 x 50 ml - Type III amber glass bottle of 50 ml closed with tamper-evident HDPE screw caps with ring and colourless LDPE syringe insert. Each bottle is individually packed in a cardboard box

10 x 10 ml - Type III amber glass bottles of 10 ml closed with tamper-evident HDPE screw caps with ring and colourless LDPE syringe insert. Each bottle is individually packed in a cardboard box

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:
Avimedical B.V.

Marketing authorisation date:
25/02/2020

Manufacturing sites for batch release:
Floris Veterinaire Producten B.V.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V554195

Date of authorisation status change:
25/02/2020

Reference member state:
Netherlands

Procedure number:
NL/V/0272/001

Concerned member states:
Belgium

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.

Labelling

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