ZORABEL 25 mg/ml oral solution chickens and turkeys

Authorised

Toltrazuril

Product identification

Medicine name:

ZORABEL 25 mg/ml oral solution chickens and turkeys ZORABEL 25 MG/ML SOLUTION POUR ADMINISTRATION DANS L'EAU DE BOISSON POUR POULETS ET DINDES

Active substance:

Toltrazuril

Target species:

Turkey

Chicken (for reproduction)

Chicken (pullet)

Route of administration:

In drinking water use

Product details

Active substance and strength:

Toltrazuril

25.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Withdrawal period by route of administration: In drinking water use:

- Turkey
 - Meat and offal. 16 day
- Chicken (for reproduction)
 - Meat and offal. 18 day
 - Eggs. no withdrawal period

Not for use in birds producing or intended to produce eggs for human consumption. Do not use within 4 weeks of the start of the laying period.

- . Chicken (pullet)
 - Meat and offal. 18 day
 - Eggs. no withdrawal period

Not for use in birds producing or intended to produce eggs for human consumption. Do not use within 4 weeks of the start of the laying period.

Anatomical therapeutic chemical veterinary (ATCvet) codes: QP51AJ01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

bottle of 5 L

bottle of 1 L

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:

Vetpharma Animal Health S.L.

Marketing authorisation date:

21/10/2013

Manufacturing sites for batch release:

Laboratorios Karizoo S.A.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/4557026 3/2013

Date of authorisation status change:

18/04/2019

Reference member state:

Spain

Procedure number:

ES/V/0199/001

Concerned member states:

France Germany Greece Poland Portugal United Kingdom (Northern Ireland)

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

Summary of Product Characteristics English (PDF) Published on: 6/04/2023 Download Package Leaflet Labelling

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