

Toltramax 50 mg/ml oral suspension for pigs

Authorised

- Toltrazuril

Product identification

Medicine name:

Toltramax 50 mg/ml oral suspension for pigs

Active substance:

Toltrazuril

Target species:

Pig

Route of administration:

Oral use

Product details

Active substance and strength:

Toltrazuril

50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral suspension

Withdrawal period by route of administration:**Oral use:**

-

Pig

- Meat and offal. 77 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP51AJ01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

White high density polyethylene bottles containing 250 or 1000 ml of suspension with a white high density polyethylene screw cap.

White high density polyethylene bottles containing 250 or 1000 ml of suspension with a white high density polyethylene screw cap.

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:

Lavet Kft.

Marketing authorisation date:

19/03/2012

Manufacturing sites for batch release:

Lavet Kft.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/6510112 2/2012

Date of authorisation status change:

18/11/2016

Reference member state:

Hungary

Procedure number:

HU/V/0114/001

Concerned member states:

Austria Belgium Cyprus Czechia Denmark France Germany Greece Italy
Lithuania Netherlands Poland Portugal Romania Spain
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.

Combined File of all Documents