# 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:**
Denmark

**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:**
17/09/2012

**Manufacturing sites for batch release:**
Lavet Kft.

**Responsible authority:**
Danish Medicines Agency

**Authorisation number:**
49844

**Date of authorisation status change:**
17/09/2012

**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

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Combined File of all Documents