

Toltarox 50 mg/ml oral suspension for cattle

Authorised

- Toltrazuril

Product identification

Medicine name:

Toltarox 50 mg/ml oral suspension for cattle

Active substance:

Toltrazuril

Target species:

Cattle

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:**

-

Cattle

- Meat and offal. 63 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP51AJ01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovenia

Package description:

Bottle (HDPE), closure (HDPE), sealing liner (LDPE): 1000 ml of oral suspension

Bottle (HDPE), closure (HDPE), sealing liner (LDPE): 250 ml of oral suspension in a 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:

KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:

24/10/2013

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:

DC/V/0335/002

Date of authorisation status change:

24/10/2013

Reference member state:

Ireland

Procedure number:

IE/V/0247/002

Concerned member states:

Romania Slovenia

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 10/05/2026

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Package Leaflet

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Labelling

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Combined File of all Documents