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: 3/05/2024
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Package Leaflet

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Labelling

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