Ridacox 50 mg/ml oral suspension for cattle and sheep

Authorised

- Toltrazuril

Product identification

Medicine name:
Ridacox 50 mg/ml oral suspension for cattle and sheep

Active substance:
Toltrazuril

Target species:
Cattle
Sheep

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
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

- **Sheep**
  - Meat and offal. 42 day
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP51BC01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Ireland

Package description:
Bottle (HDPE), closure (HDPE), sealing liner (LDPE): 250 ml of oral suspension in a box.
Bottle (HDPE), closure (HDPE), sealing liner (LDPE): 1000 ml of oral suspension in a box.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 10(1) of Directive No 2001/83/EC)

Marketing authorisation holder:
J & M Veterinary Services Limited

Marketing authorisation date:
17/04/2015

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10954/012/001

Date of authorisation status change:
17/04/2015

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

Documents

Summary of Product Characteristics

ie-puar-np-600000064498-ridacox-50-mgml-oral-suspension-for-cattle-and-she-en.pdf