

Bovicox 50 mg/ml oral suspension for cattle and sheep

Authorised

- Toltrazuril

Product identification

Medicine name:

Bovicox 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

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 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

J & M Veterinary Services Limited

Marketing authorisation date:

20/09/2013

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10954/011/001

Date of authorisation status change:

20/09/2013

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

Documents

Summary of Product Characteristics

ie-puar-np-600000064516-bovicox-50-mgml-oral-suspension-for-cattle-and-she-en.pdf