

Baycox Multi 50 mg/ml oral suspension for Cattle, Pigs and Sheep

Authorised

- Toltrazuril

Product identification

Medicine name:

Baycox Multi 50 mg/ml oral suspension for Cattle, Pigs and Sheep

Baycox Multi 50 mg/ml πόσιμο εναιώρημα για βοοειδή, χοίρους και πρόβατα

Active substance:

Toltrazuril

Target species:

Cattle

Sheep

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:

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Cattle

- Meat and offal. 63 day

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Sheep

- Meat and offal. 42 day

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Pig

- Meat and offal. 77 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP51BC01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Cyprus

Available in:

Cyprus

Package description:

100 ml high density polyethylene bottles closed with polypropylene screw caps.

250 ml high density polyethylene bottles closed with polypropylene screw caps.

1000 ml high density polyethylene bottles closed with polypropylene screw caps.

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:

Elanco Animal Health GmbH

Marketing authorisation date:

27/06/2017

Manufacturing sites for batch release:

KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

CY00619V

Date of authorisation status change:

7/11/2021

Reference member state:

Ireland

Procedure number:

IE/V/0360/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland
France Germany Greece Hungary Iceland Italy Latvia Lithuania
Luxembourg Netherlands Norway Poland Portugal Romania Slovakia
Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

[Summary of Product Characteristics](#)

[Combined File of all Documents](#)

English (PDF)

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