Chanox Multi 50 mg/ml oral suspension for Piglets, Calves and Lambs

Authorised

- Toltrazuril

Product identification

Medicine name:
Chanox Multi 50 mg/ml oral suspension for Piglets, Calves and Lambs

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:

- **Cattle**
  - Meat and offal. 63 day
- **Sheep**
  - Meat and offal. 42 day
- **Pig**
  - Meat and offal. 77 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP51AJ01

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

Authorisation status:
Valid

Authorised in:
Cyprus

Package description:
High density polyethylene flexi-pack bottles containing 5 L with a polypropylene screw cap closure.
High density polyethylene flexi-pack bottles containing 1 L with a polypropylene screw cap closure.
High density polyethylene bottles containing 250 ml with a high density polyethylene screw cap closure.
High density polyethylene bottles containing 100 ml with a high density polyethylene screw cap closure.

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:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
3/12/2017

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00650V

Date of authorisation status change:
3/12/2017

Reference member state:
Ireland

Procedure number:
IE/V/0628/001

Concerned member states:
Austria Belgium Cyprus Czechia Finland France Germany Greece Hungary Italy Netherlands Norway Poland Portugal Romania Slovakia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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