

Chanil 34 mg/ml oral suspension for cattle

Authorised

- Oxyclozanide

Product identification

Medicine name:

Chanil 34 mg/ml oral suspension for cattle
RUMENIL 34 mg/ml suspensie orala pentru bovine

Active substance:

Oxyclozanide

Target species:

Cattle

Route of administration:

Oral use

Product details

Active substance and strength:

Oxyclozanide
34.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral suspension

Withdrawal period by route of administration:

Oral use:**• Cattle**

- Meat and offal. 13 day
 - Milk. 108 hour
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AG06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Romania

Package description:

10 L: High Density Polyethylene (HDPE) container with a HDPE cap and an aluminium foil seal. The product can be marketed with or without an outer carton.

5L:White High Density Polyethylene HDPE flexi containers with a Polypropylene cap and a PVDC seal The product can be marketed with or without an outer carton.

2.5L:White High Density Polyethylene HDPE flexi containers with a Polypropylene cap and a PVDC seal The product can be marketed with or without an outer carton.

1L:White High Density Polyethylene HDPE flexi containers with a Polypropylene cap and a PVDC seal. The product can be marketed with or without an outer carton.

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:

19/11/2017

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

210202

Date of authorisation status change:

14/12/2021

Reference member state:

Ireland

Procedure number:

IE/V/0368/001

Concerned member states:

Austria Belgium France Netherlands Romania

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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