

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 SUSPENSION BUVABLE POUR BOVINS

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:**

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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:

France

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:

15/12/2017

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/5710098 8/2017

Date of authorisation status change:

12/03/2021

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: 4/05/2025

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Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents