

# Chanal 34 mg/ml oral suspension for cattle

Authorised

- Oxyclozanide

## Product identification

**Medicine name:**

Chanal 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:**

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**Cattle**

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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QP52AG06

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Romania

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Chanelle Pharmaceuticals Manufacturing Limited

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**Marketing authorisation date:**

19/11/2017

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**Manufacturing sites for batch release:**

Chanelle Pharmaceuticals Manufacturing Limited

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**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

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**Authorisation number:**

210202

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**Date of authorisation status change:**

12/05/2025

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0368/001

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**Concerned member states:**

Austria Belgium France Netherlands Romania

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

## Combined File of all Documents

English (PDF)

Published on: 4/05/2025

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