

# Rumenil 34 mg/ml oral suspension for cattle

Not  
authorised

- Oxyclozanide

## Product identification

**Medicine name:**

Rumenil 34 mg/ml oral suspension for cattle

Rumenil 34 mg/ml перорална суспензия за говеда

**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:**

QP52AG05

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

Bulgaria

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**Package description:**

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.

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.

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.

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.

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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:**

16/09/2016

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**Manufacturing sites for batch release:**

Chanelle Pharmaceuticals Manufacturing Limited

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**Responsible authority:**

Bulgarian Food Safety Authority

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**Authorisation number:**

0022-2679

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**Date of authorisation status change:**

28/08/2023

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0369/001

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## Documents

Summary of Product Characteristics

English (PDF)

Published on: 4/05/2025

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Package Leaflet and Labelling

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Combined File of all Documents

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