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 voor oraal gebruik
Rumenil 34 mg/ml Suspension buvable
Rumenil 34 mg/ml Suspension zum Einnehmen

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:
Belgium

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 sealThe 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 sealThe 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:
10/09/2018

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
10/09/2018

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

Download

Package Leaflet

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

Combined File of all Documents

Package Leaflet and Labelling

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