Rumenil 34 mg/ml oral suspension for cattle

Authorised

- Oxyclozanide

Product identification

Medicine name:
Rumenil 34 mg/ml oral suspension for cattle

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:

QP52AG05

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Package description:

1L: 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.
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.
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.

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:
9/09/2016

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
3465 ESP

Date of authorisation status change:
9/09/2016

Reference member state:
Ireland

Procedure number:
IE/V/0369/001

Concerned member states:
Bulgaria Hungary Italy Spain United Kingdom (Northern Ireland)

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

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Labelling

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Combined File of all Documents