Oxyfluke 34 mg/ml Oral Suspension for Cattle and Sheep

Authorised

- Oxyclozanide

Product identification

Medicine name:
Oxyfluke 34 mg/ml Oral Suspension for Cattle and Sheep

Active substance:
Oxyclozanide

Target species:
Cattle
Sheep

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
- **Sheep**
  - Meat and offal. 14 day
  - Milk. 7 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AG06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Slovenia

Package description:

White high density polyethylene backpacks (1 L) closed with white polypropylene screw caps. 1 x 1 L in a cardboard box

White high density polyethylene backpacks (2.5 L) closed with white polypropylene screw caps. 1 x 2.5 L in a cardboard box

White high density polyethylene backpacks (5 L) closed with white polypropylene screw caps. 1 x 5 L in a cardboard box

White high density polyethylene backpacks (5 L) closed with white polypropylene screw caps. 2 x 5 L in a cardboard box

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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:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
18/12/2017

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0596/001

Date of authorisation status change:
18/12/2017

Reference member state:
Ireland

Procedure number:
IE/V/0601/001

Concerned member states:
Belgium Bulgaria Croatia Hungary Luxembourg Netherlands Portugal Slovenia 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: 3/05/2024
Download

Package Leaflet

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Labelling

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