

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:**

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Cattle

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

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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

Authorisation status:

Valid

Authorised in:

Luxembourg

Package description:

White high density polyethylene backpacks (5 L) closed with white polypropylene screw caps. 2 x 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 (2.5 L) closed with white polypropylene screw caps. 1 x 2.5 L in a cardboard box

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

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:

15/11/2017

Manufacturing sites for batch release:

Norbrook Laboratories Limited

Norbrook Manufacturing Limited

Responsible authority:

Ministry Of Health And Social Security

Authorisation number:

IE/V/0601/001

Date of authorisation status change:

11/04/2022

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