

Distocur 34 mg/ml Oral suspension for cattle and sheep

Authorised

- Oxyclozanide

Product identification

Medicine name:

Distocur 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. 5 day

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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:

Ireland

Available in:

Ireland

Package description:

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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:

Dopharma Research B.V.

Marketing authorisation date:

13/01/2017

Manufacturing sites for batch release:

Dopharma France

Dopharma B.V.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10791/013/001

Date of authorisation status change:

13/01/2017

Reference member state:

France

Procedure number:

FR/V/0312/001

Concerned member states:

Austria Belgium Croatia Denmark Germany Hungary Ireland Italy
Luxembourg Netherlands Norway Poland Portugal Romania Slovenia
Sweden United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Combined File of all Documents

eu-puar-frv0312001-mr-rpe515-en.pdf