

Flordofen 300 mg/ml solution for injection for cattle and pigs

Authorised

- Florfenicol

Product identification

Medicine name:

Flordofen 300 mg/ml solution for injection for cattle and pigs

Active substance:

Florfenicol

Target species:

Pig (for fattening)

Cattle

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Florfenicol

300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Pig (for fattening)

- Meat and offal. 18 day

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Cattle

- Meat and offal. 30 day

Milk: Not authorised for use in animals producing milk for human consumption, including during the dry period.

Subcutaneous use:

-

Cattle

- Meat and offal. 44 day

Milk: Not authorised for use in animals producing milk for human consumption, including during the dry period.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01BA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Package description:

Brown-coloured type II glass vial of 250 ml, closed by a type I bromobutyl stopper and sealed by an aluminium cap with centre hole.

Colourless type II glass vial of 100 ml, closed by a type I bromobutyl stopper and sealed by an aluminium cap with centre hole.

Colourless type II glass vial of 50 ml, closed by a type I bromobutyl stopper and sealed by an aluminium cap with centre hole.

Polypropylene vial of 250 ml, closed with bromobutyl stopper secured with flip off aluminium collar.

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:

4/07/2019

Manufacturing sites for batch release:

Dopharma B.V.

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-2903

Date of authorisation status change:

4/07/2019

Reference member state:

Portugal

Procedure number:

PT/V/0112/001

Concerned member states:

Austria Belgium Bulgaria Croatia Czechia Denmark Estonia France
Germany Greece Hungary Ireland Italy Latvia Lithuania Netherlands Poland
Romania Slovakia 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

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

Package Leaflet

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

Combined File of all Documents