

# FLORON 300 MG/ML SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Authorised

- Florfenicol

## Product identification

**Medicine name:**

FLORON 300 MG/ML SOLUTION FOR INJECTION FOR CATTLE AND PIGS

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**Active substance:**

Florfenicol

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**Target species:**

Cattle

Pig

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

**Active substance and strength:**

Florfenicol

300.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Cattle**

- Meat and offal. 30 day dosage 20 mg/kg bodyweight, twice
- Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

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**Pig**

- Meat and offal. 18 day

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 44 day dosage 40 mg/kg bodyweight, once
- Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01BA90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

France

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**Package description:**

Cardboard box containing one Type I amber glass bottle of 50 ml  
Cardboard box containing one Type I amber glass bottle of 250 ml  
Cardboard box containing one Type I amber glass bottle of 100 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

KRKA tovarna zdravil d.d. Novo mesto

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**Marketing authorisation date:**

5/01/2015

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**Manufacturing sites for batch release:**

TAD Pharma GmbH

KRKA tovarna zdravil d.d. Novo mesto

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/1005669 1/2014

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**Date of authorisation status change:**

16/12/2019

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**Reference member state:**

France

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**Procedure number:**

FR/V/0280/001

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**Concerned member states:**

Belgium Germany Italy Netherlands Portugal Spain

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://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.

Combined File of all Documents

Package Leaflet and Labelling

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