

# RESFLOR 300/16.5 MG/ML SOLUTION FOR INJECTION FOR CATTLE

Authorised

- Florfenicol
- Flunixin meglumine

## Product identification

**Medicine name:**

RESFLOR 300/16.5 MG/ML SOLUTION FOR INJECTION FOR CATTLE

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**Active substance:**

Florfenicol

Flunixin meglumine

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**Target species:**

Cattle

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Florfenicol

300.00 milligram(s) / 1.00 millilitre(s)

Flunixin meglumine  
27.40 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:**

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 46 day
- Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption. Do not use in pregnant animals which are intended to produce milk for human consumption within 2 months of expected parturition.

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

QJ01BA99

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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:**

Norway

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**Available in:**

Norway

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**Package description:**

Carton box containing 100 mL vial  
Carton box containing 250 mL vial

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

24/08/2007

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**Manufacturing sites for batch release:**

Vet Pharma Friesoythe GmbH

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**Responsible authority:**

Norwegian Medical Products Agency

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**Authorisation number:**

07-4761

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**Date of authorisation status change:**

6/01/2011

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**Reference member state:**

France

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**Procedure number:**

FR/V/0167/001

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**Concerned member states:**

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland  
Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg  
Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain  
United Kingdom (Northern Ireland)

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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.

### Package Leaflet

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