

# MYCOFLOR 300 mg/ml solution for injection for cattle and pigs

Authorised

- Florfenicol

## Product identification

**Medicine name:**

MYCOFLOR 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

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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. 34 day

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**Pig**

- Meat and offal. 18 day

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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:**

Portugal

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**Available in:**

Portugal

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**Package description:**

Vial size: 100 ml Colourless Type II glass vial closed with bromobutyl rubber closure.

Vials are individually packed in carton box.

Vial size: 250 ml Colourless Type II glass vial closed with bromobutyl rubber closure.

Vials are individually packed in carton box.

Vial size: 100 ml An aluminium cap or polypropylene vial closed with bromobutyl rubber closure and an aluminium cap. Vials are individually packed in carton box.

Vial size: 250 ml An aluminium cap or polypropylene vial closed with bromobutyl rubber closure and an aluminium cap. Vials are individually packed in carton box.

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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:**

S P Veterinaria S.A.

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**Marketing authorisation date:**

19/12/2011

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**Manufacturing sites for batch release:**

KELA Kempisch Laboratorium Kela Laboratoria  
S P Veterinaria S.A.

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**Responsible authority:**

Directorate General For Food And Veterinary

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**Authorisation number:**

390/01/11DFVPT

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**Date of authorisation status change:**

29/08/2024

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0486/001

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**Concerned member states:**

Belgium Bulgaria Cyprus Denmark France Germany Greece Hungary Italy  
Poland Portugal Romania 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

Combined File of all Documents

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