

# Norfenicol 300 mg/ml Solution for Injection for Cattle and Pigs

Not authorised

- Florfenicol

## Product identification

**Medicine name:**

Norfenicol 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. 39 day

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**Pig**

- Meat and offal. 22 day

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 44 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:**

Surrendered

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**Authorised in:**

Belgium

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**Package description:**

500 ml clear type I glass vial with bromobutyl rubber bung and aluminium seal accompanied by a protective sleeve.

250 ml clear type I glass vial with bromobutyl rubber bung and aluminium seal accompanied by a protective sleeve.

100 ml clear type I glass vial with bromobutyl rubber bung and aluminium seal accompanied by a protective sleeve.

50 ml clear type I glass vial with bromobutyl rubber bung and aluminium seal presented in a cardboard box.

50 ml HDPE plastic vial with bromobutyl rubber bung and aluminium seal presented in a cardboard box.

100 ml HDPE plastic vial with bromobutyl rubber bung and aluminium seal presented in a cardboard box.

250 ml HDPE plastic vial with bromobutyl rubber bung and aluminium seal presented in a cardboard box.

500 ml HDPE plastic vial with bromobutyl rubber bung and aluminium seal presented in a cardboard 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:**

Norbrook Laboratories (Ireland) Limited

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### **Marketing authorisation date:**

28/09/2012

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### **Manufacturing sites for batch release:**

Norbrook Laboratories Limited  
Norbrook Manufacturing Limited

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### **Responsible authority:**

Federal Agency For Medicines And Health Products

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### **Authorisation number:**

This information is not available for this product.

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### **Date of authorisation status change:**

21/03/2025

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### **Reference member state:**

Ireland

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**Procedure number:**IE/V/0282/001

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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

English (PDF)

Published on: 16/01/2026

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### Package Leaflet

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