

# Nuflor Minidose 450 mg/ml Solution for Injection for Cattle

Authorised

- Florfenicol

## Product identification

**Medicine name:**

Nuflor Minidose 450 mg/ml solution for injection for cattle

Nuflor Minidose 450 mg/ml Solution for Injection for Cattle

**Active substance:**

Florfenicol

**Target species:**

Cattle

**Route of administration:**

Subcutaneous use

Intramuscular use

## Product details

**Active substance and strength:**

Florfenicol

450.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Cattle**

- Milk. no withdrawal period

Not permitted for use in lactating animals producing milk for human consumption.

- Meat and offal. 64 day

**Intramuscular use:**

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**Cattle**

- Milk. no withdrawal period

Not permitted for use in lactating animals producing milk for human consumption.

- Meat and offal. 37 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:**

United Kingdom (Northern Ireland)

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**Package description:**

(ID3) 250 millilitre(s): unspecified outer container with 1 Vial (Glass) with 250 millilitre(s), closed with Stopper (bromobutyl rubber`)

(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (Glass) with 100 millilitre(s), closed with Stopper (bromobutyl rubber`)

(ID1) 50 millilitre(s): unspecified outer container with 1 Vial (Glass) with 50 millilitre(s), closed with Stopper (bromobutyl rubber`)

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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:**

Intervet International B.V.

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**Marketing authorisation date:**

3/10/2008

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**Manufacturing sites for batch release:**

Intervet International GmbH

Trirx Segre

Vet Pharma Friesoythe GmbH

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**Responsible authority:**

The Veterinary Medicines Directorate

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**Authorisation number:**

Vm 06376/3048

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**Date of authorisation status change:**

28/04/2024

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0122/001

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**Concerned member states:**

Belgium Bulgaria Cyprus Czechia Denmark Finland France Greece Hungary

Ireland Italy Luxembourg Netherlands Portugal Romania 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

Combined File of all Documents