

# 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

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**Active substance:**

Florfenicol

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**Target species:**

Cattle

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**Route of administration:**

Subcutaneous use  
Intramuscular use

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## Product details

**Active substance and strength:**

Florfenicol  
450.00 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:****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:**

France

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**Available in:**

France

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**Package description:**

(ID1) 50 millilitre(s): unspecified outer container with 1 Vial (glass) with 50 millilitre(s), closed with Stopfen (bromobutyl rubber)

(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (glass) with 100 millilitre(s), closed with Stopfen (bromobutyl rubber)

(ID3) 250 millilitre(s): unspecified outer container with 1 Vial (glass) with 250 millilitre(s), closed with Stopfen (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:**

17/09/2008

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**Manufacturing sites for batch release:**

Intervet International GmbH

Trirx Segre

Vet Pharma Friesoythe GmbH

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/7051353 4/2008

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**Date of authorisation status change:**

17/09/2013

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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

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

Package Leaflet and Labelling

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

Combined File of all Documents

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