

# FLORFENICEN 300 mg/ml solution for injection for cattle, sheep and pig

Authorised

- Florfenicol
- Florfenicol

## Product identification

### Medicine name:

FLORFENICEN 300 mg/ml solution for injection for cattle, sheep and pig

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### Active substance:

Florfenicol

Florfenicol

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### Target species:

Cattle

Sheep

Pig

Cattle

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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)

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. 30 day por via IM (a 20mg/kg de peso corporal, duas vezes)

- Meat and offal. 44 day por via SC (a 40 mg/kg de peso corporal, uma vez)

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#### Sheep

- Meat and offal. 39 day

Não administrar a fêmeas produtoras de leite destinado ao consumo humano, incluindo animais gestantes com intenção de produção de leite para consumo humano

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#### Pig

- Meat and offal. 18 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:**

Valid

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**Authorised in:**

Portugal

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**Package description:**

250 ml polypropylene vials closed with a rubber-butyl septum and an aluminium capsule with blue Flip-Off sealing.

100 ml polypropylene vials closed with a rubber-butyl septum and an aluminium capsule with blue Flip-Off sealing.

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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:**

Cenavisa S.L.

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**Marketing authorisation date:**

6/08/2012

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**Manufacturing sites for batch release:**

Cenavisa S.L.

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**Responsible authority:**

Directorate General For Food And Veterinary

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**Authorisation number:**

586/01/12DFVPT

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**Date of authorisation status change:**

5/04/2022

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**Reference member state:**

Portugal

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**Procedure number:**

PT/V/0106/001

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**Concerned member states:**

Spain

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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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