

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

Active substance:

Florfenicol

Florfenicol

Target species:

Cattle

Sheep

Pig

Cattle

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Florfenicol

300.00 milligram(s) / 1.00 millilitre(s)

Florfenicol

300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

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:

-

Cattle

- Meat and offal. 44 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01BA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

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.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Cenavisa S.L.

Marketing authorisation date:

7/09/2012

Manufacturing sites for batch release:

Cenavisa S.L.

Responsible authority:

Spanish Agency For Medicines And Medical Devices

Authorisation number:

2621 ESP

Date of authorisation status change:

7/09/2012

Reference member state:

Portugal

Procedure number:

PT/V/0106/001

Concerned member states:

Spain

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Labelling

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Summary of Product Characteristics

English (PDF)

Published on: 25/11/2021

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Package Leaflet

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