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
CENFLOR 300 mg/ml solução injetável para bovinos, ovinos e suínos

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)
- **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

- **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:
Portugal

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:
6/08/2012

Manufacturing sites for batch release:
Cenavisa S.L.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:

586/01/12DFVPT

Date of authorisation status change:
5/04/2022

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

Summary of Product Characteristics

Combined File of all Documents

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