PULMOVALL 300 mg/ml solution for injection for cattle, sheep and pigs.

Authorised

- Florfenicol

Product identification

Medicine name:
PULMOVALL 300 mg/ml solution for injection for cattle, sheep and pigs.

Active substance:
Florfenicol

Target species:
Cattle
Sheep
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
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. no withdrawal period

Meat and offal: Intramuscular use (20 mg / kg bw, twice): 30 days / Subcutaneous use (40 mg / kg bw, once): 44 days

- **Sheep**
  - Meat and offal. 39 day

- **Pig**
  - Meat and offal. 18 day

- **Cattle**
  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption including pregnant animals intended to produce milk for human consumption.

- **Sheep**
  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption including pregnant animals intended to produce milk for human consumption.

Subcutaneous use:

- **Cattle**

- Meat and offal. no withdrawal period

Meat and offal: Intramuscular use (20 mg / kg bw, twice): 30 days / Subcutaneous use (40 mg / kg bw, once): 44 days

- **Cattle**
  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption including pregnant animals intended to produce milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01BA90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Portugal

Available in:
Portugal

Package description:
cardboard box containing 15 vials of 250 ml
cardboard box containing 10 vials of 100 ml
cardboard box containing 1 vial of 250 ml
cardboard box containing 1 vial of 100 ml

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:
Mevet S.A.

Marketing authorisation date:
23/02/2021

Manufacturing sites for batch release:
Mevet S.A.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
1407/01/21DFVPT

Date of authorisation status change:
4/04/2024

Reference member state:
Spain

Procedure number:
ES/V/0376/001

Concerned member states:
Poland Portugal Romania

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

Documents

Combined File of all Documents

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

Summary of Product Characteristics

Package Leaflet

Labelling

eu-PUAR-pulmovall-300-mg-ml-solution-for-injection-for-cattle--sheep-and-pigs.-en.pdf