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

Authorisation status:
Valid

Authorised in:
Romania

Available in:
Romania

Package description:
cardboard box containing 1 vial of 100 ml
cardboard box containing 1 vial of 250 ml
cardboard box containing 10 vials of 100 ml
cardboard box containing 15 vials of 250 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:
10/03/2021

Manufacturing sites for batch release:
Mevet S.A.

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
260042

Date of authorisation status change:
9/03/2026

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

Summary of Product Characteristics

Package Leaflet

Labelling

Combined File of all Documents

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