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

Authorised

- Florfenicol

Product identification

Medicine name:
Pulmovall 300 mg/ml Roztwór do wstrzykiwań
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:
Poland

Package description:
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish

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:
22/03/2021

Manufacturing sites for batch release:
Mevet S.A.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3086

Date of authorisation status change:
22/03/2021

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

Labelling

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Package Leaflet

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

Summary of Product Characteristics

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