

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

Authorised

- Florfenicol

Product identification

Medicine name:

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

Active substance:

Florfenicol

Target species:

Cattle

Pig

Sheep

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:

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Cattle

- Meat and offal. 30 day
- 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.

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Pig

- Meat and offal. 18 day

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Sheep

- Meat and offal. 39 day
- 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.

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Cattle

- Meat and offal. 30 day
- 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.

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Sheep

- Meat and offal. 39 day

- 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:

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Cattle

- Meat and offal. 44 day
- 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.

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Cattle

- Meat and offal. 44 day
- 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:

Portugal

Available in:

Portugal

Package description:

box containing 6 vials of 250 ml

box containing 6 vials of 100 ml
box containing 1 vial of 250 ml
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:

Laboratorios Syva S.A.

Marketing authorisation date:

5/01/2021

Manufacturing sites for batch release:

Laboratorios Syva S.A.

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

1393/01/21DFVPT

Date of authorisation status change:

8/02/2023

Reference member state:

Spain

Procedure number:

ES/V/0377/001

Concerned member states:

Belgium Denmark France Germany Greece Hungary Ireland Italy Lithuania
Netherlands 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

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