

Mycoflor, 300 mg/ml Solution for Injection for Cattle and Pigs

Authorised

- Florfenicol

Product identification

Medicine name:

Mycoflor, 300 mg/ml Solution for Injection for Cattle and Pigs

Active substance:

Florfenicol

Target species:

Cattle

Pig

Route of administration:

Intramuscular 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. 34 day

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Pig

- Meat and offal. 18 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01BA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Package description:

Vial size: 100 ml An aluminium cap or polypropylene vial closed with bromobutyl rubber closure and an aluminium cap. Vials are individually packed in carton box.

Vial size: 250 ml An aluminium cap or polypropylene vial closed with bromobutyl rubber closure and an aluminium cap. Vials are individually packed in carton box.

Vial size: 100 ml Colourless Type II glass vial closed with bromobutyl rubber closure. Vials are individually packed in carton box.

Vial size: 250 ml Colourless Type II glass vial closed with bromobutyl rubber closure. Vials are individually packed in carton box.

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:

S P Veterinaria S.A.

Marketing authorisation date:

21/12/2011

Manufacturing sites for batch release:

KELA Kempisch Laboratorium Kela Laboratoria
S P Veterinaria S.A.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 36967/3004

Date of authorisation status change:

6/07/2024

Reference member state:

Ireland

Procedure number:

IE/V/0486/001

Concerned member states:

Belgium Bulgaria Cyprus Denmark France Germany Greece Hungary Italy
Poland Portugal Romania Spain United Kingdom (Northern Ireland)

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