FLORKEM 300 MG/ML SOLUTION FOR INJECTION FOR CATTLE AND PIGS

Authorised

- Florfenicol

Product identification

Medicine name:
FLORKEM 300 MG/ML SOLUTION FOR INJECTION FOR CATTLE AND PIGS
FLORKEM 300 mg/ml SOLUCION INYECTABLE PARA BOVINO Y PORCINO

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:

- **Cattle**
  - Meat and offal. 37 day
  - Milk. no withdrawal period

Milk: Not permitted for use in lactating animals producing milk for human consumption.

- **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:

Spain

Available in:

Spain

Package description:

Box containing one glass vial of 20 ml
Box containing one plastic vial of 500 ml
Box containing one plastic vial of 250 ml
Box containing one plastic vial of 100 ml
Box containing one plastic vial of 50 ml
Box containing one glass vial of 500 ml

Box containing one glass vial of 250 ml
Box containing one glass vial of 100 ml
Box containing one glass vial of 50 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:
Ceva Salud Animal S.A.

Marketing authorisation date:
10/07/2009

Manufacturing sites for batch release:
CEVA Santé Animale

Responsible authority:
Spanish Agency For Medicines And Medical Devices

Authorisation number:
2051 ESP

Date of authorisation status change:
10/07/2009

Reference member state:
France

Procedure number:
FR/V/0197/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands

Poland Portugal Romania Slovakia 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

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

Package Leaflet

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

Labelling

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