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

Authorised

- Florfenicol

Product identification

Medicine name:
FLORON 300 MG/ML SOLUTION FOR INJECTION FOR CATTLE AND PIGS
Floron 300 mg/ml Injektionslösung für Rinder und Schweine

Active substance:
Florfenicol

Target species:
Cattle
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. 30 day — dosage 20 mg/kg bodyweight, twice
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

- **Pig**
  - Meat and offal. 18 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 44 day — dosage 40 mg/kg bodyweight, once
  - Milk. no withdrawal period

Not authorised for use in animals producing 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:
Germany

Package description:

Cardboard box containing one Type I amber glass bottle of 100 ml
Cardboard box containing one Type I amber glass bottle of 250 ml
Cardboard box containing one Type I amber glass bottle 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:
TAD Pharma GmbH

Marketing authorisation date:
20/01/2015

Manufacturing sites for batch release:
TAD Pharma GmbH
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402120.00.00

Date of authorisation status change:
20/01/2020

Reference member state:
France

Procedure number:
FR/V/0280/001

Concerned member states:
Belgium Germany Italy Netherlands Portugal Spain

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

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