RESFLOR 300/16.5 MG/ML SOLUTION FOR INJECTION FOR CATTLE

Authorised

- Florfenicol
- Flunixin meglumine

Product identification

Medicine name:
RESFLOR 300/16.5 MG/ML SOLUTION FOR INJECTION FOR CATTLE
Resflor vet. 300 mg/ml / 16,5 mg/ml injeksjonsvæske, oppløsning til storfe

Active substance:
Florfenicol
Flunixin meglumine

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Florfenicol

300.00 milligram(s) / 1.00 millilitre(s)

Flunixin meglumine
27.40 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Cattle**
  - Meat and offal. 46 day
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption. Do not use in pregnant animals which are intended to produce milk for human consumption within 2 months of expected parturition.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01BA99

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Norway

Package description:
Carton box containing 100 mL vial
Carton box containing 250 mL vial

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
24/08/2007

Manufacturing sites for batch release:
Vet Pharma Friesoythe GmbH

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
07-4761

Date of authorisation status change:
6/01/2011

Reference member state:
France

Procedure number:
FR/V/0167/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia 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.