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

Not authorised

- Florfenicol
- Flunixin meglumine

Product identification

Medicine name:
RESFLOR 300/16.5 MG/ML SOLUTION FOR INJECTION FOR CATTLE

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:
Surrendered

Authorised in:
Finland

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:
11/07/2010

Manufacturing sites for batch release:
Vet Pharma Friesoythe GmbH

Responsible authority:
Finnish Medicines Agency

Authorisation number:
23260

Date of authorisation status change:
13/03/2023

Reference member state:
France

Procedure number:
FR/V/0167/001

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

Documents

Combined File of all Documents