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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