Norfenicol 300 mg/ml Solution for Injection for Cattle and Pigs

Authorised

- Florfenicol

Product identification

Medicine name:
Norfenicol 300 mg/ml Solution for Injection for Cattle and Pigs
Norfenicol 300 mg/ml injektionsvæske, opløsning

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. 39 day

- **Pig**

  - Meat and offal. 22 day

Subcutaneous use:

- **Cattle**

  - Meat and offal. 44 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01BA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Denmark

Package description:

500 ml clear type I glass vial with bromobutyl rubber bung and aluminium seal presented in a cardboard box or a protective sleeve.
250 ml clear type I glass vial with bromobutyl rubber bung and aluminium presented in a cardboard box or a protective sleeve.
100 ml clear type I glass vial with bromobutyl rubber bung and aluminium seal presented in a cardboard box or a protective sleeve.
50 ml clear type I glass vial with bromobutyl rubber bung and aluminium seal presented in a cardboard box.
50 ml HDPE plastic vial with bromobutyl rubber bung and aluminium seal presented in a cardboard box.
100 ml HDPE plastic vial with bromobutyl rubber bung and aluminium seal presented in a cardboard box or a protective sleeve.
250 ml HDPE plastic vial with bromobutyl rubber bung and aluminium seal presented in a cardboard box or a protective sleeve.

500 ml HDPE plastic vial with bromobutyl rubber bung and aluminium seal presented in a cardboard box or a protective sleeve.

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:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
This information is not available for this product.

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Danish Medicines Agency

Authorisation number:
48668

Date of authorisation status change:
21/05/2012

Reference member state:
Ireland

Procedure number:
IE/V/0282/001

Concerned member states:
Belgium Czechia Denmark Germany Greece Hungary Italy Luxembourg Netherlands 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

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