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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

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

Authorisation status:

Valid

Authorised in:

Slovenia

Package description:

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

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

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:
2/03/2012

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

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0385/001

Date of authorisation status change:
2/03/2012

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 United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 26/07/2026
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Combined File of all Documents

Labelling

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Package Leaflet

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