

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:

United Kingdom (Northern Ireland)

Available in:

United Kingdom (Northern Ireland)

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 seal 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 Limited

Marketing authorisation date:

11/05/2012

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 02000/4316

Date of authorisation status change:

27/03/2024

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