

Nuflor 40 mg/g Premix for Medicated Feeding Stuff for Swine

Authorised

- Florfenicol

Product identification

Medicine name:

Nuflor 40 mg/g Premix for Medicated Feeding Stuff for Swine
Nuflor 40 mg/g πρόμιγμα για φαρμακώχο τροφή για χοίρους

Active substance:

Florfenicol

Target species:

Pig

Route of administration:

Oral use

Product details

Active substance and strength:

Florfenicol
40.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Premix for medicated feeding stuff

Withdrawal period by route of administration:

Oral use:

-

Pig

- Meat and offal. 14 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01BA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Cyprus

Package description:

LDPE/HDPE/paper sealed bag containing 5 kg premix.

LDPE/paper/paper/paper sealed bag containing 25 kg premix.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet Nederland B.V.

Marketing authorisation date:

16/09/2007

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

CY00134V

Date of authorisation status change:

21/06/2020

Reference member state:

Ireland

Procedure number:

IE/V/0594/001

Concerned member states:

Belgium Cyprus Czechia Finland Greece Hungary Italy Luxembourg
Netherlands Portugal 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

Combined File of all Documents

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