

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

Not authorised

- Florfenicol

Product identification

Medicine name:

Nuflor 40 mg/g Premix for Medicated Feeding Stuff for Swine
Nuflor vet. 40 mg/g esisekoite lääkerehua varten

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:

Surrendered

Authorised in:

Finland

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 International B.V.

Marketing authorisation date:

8/08/2011

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Finnish Medicines Agency

Authorisation number:

22899

Date of authorisation status change:

23/02/2023

Reference member state:

Ireland

Procedure number:

IE/V/0594/001

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

Documents

Summary of Product Characteristics

ie-puar-mr-iev0594001-nuflor-40-mgg-premix-for-medicated-feeding-stuff-f-en.pdf

Source URL: <https://medicines.health.europa.eu/veterinary/600000047285>