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