Feniveex Premix

Not authorised

- Florfenicol

Product identification

Medicine name:
Feniveex Premix

Active substance:
Florfenicol

Target species:
Pig

Route of administration:
In-feed use

Product details

Active substance and strength:
Florfenicol
2.00 gram(s) / 100.00 gram(s)

Pharmaceutical form:
Premix for medicated feeding stuff

Withdrawal period by route of administration:

In-feed use:

- **Pig**
  - Meat and offal. 15 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01BA90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Surrendered

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
S P Veterinaria S.A.

Marketing authorisation date:
2/06/2010

Manufacturing sites for batch release:
S P Veterinaria S.A.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1387

Date of authorisation status change:
26/06/2022

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

Documents

Summary of Product Characteristics

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

Package Leaflet and Labelling

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