NEO OXYVET PREMIX

Authorised

- NEOMYCIN SULFATE
- Oxytetracycline hydrochloride

Product identification

Medicine name:
НЕО ОКСИВЕТ ПРЕМИКС
NEO OXYVET PREMIX

Active substance:
NEOMYCIN SULFATE
Oxytetracycline hydrochloride

Target species:
Pig
Poultry

Route of administration:
Oral use

Product details

Active substance and strength:
NEOMYCIN SULFATE
44.00 gram(s) / 1.00 kilogram(s)
Oxytetracycline hydrochloride

44.00 gram(s) / 1.00 kilogram(s)

Pharmaceutical form:
Premix for medicated feeding stuff

Withdrawal period by route of administration:
Oral use:

- **Pig**
  - Meat and offal. 6 day
- **Poultry**
  - Meat and offal. 14 day

Не се разрешава употребата при птици, чиито яйца са предназначени за човешка консумация

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA56

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Available in:
Bulgaria

Package description:
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:
PROVET S.A.

Marketing authorisation date:
12/05/2008

Manufacturing sites for batch release:
PROVET S.A.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2041

Date of authorisation status change:
27/03/2022

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

Documents

Combined File of all Documents

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

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.