NEOMAY 500 000 IU/G POWDER FOR USE IN DRINKING WATER/MILK REPLACER

Authorised

- Neomycin

Product identification

Medicine name:
NEOMAY 500 000 IU/G POWDER FOR USE IN DRINKING WATER/MILK REPLACER

Active substance:
Neomycin

Target species:
Cattle
Pig
Poultry

Route of administration:
Oral use

Product details

Active substance and strength:
Neomycin
500000.00 international unit(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for use in drinking water/milk

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 14 day
- **Pig**
  - Meat and offal. 3 day
- **Poultry**
  - Meat and offal. 14 day
  - Eggs. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA07AA01

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

Authorisation status:
Valid

Authorised in:
Denmark

Package description:
bag of 100 g
bag of 1 kg

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Maymo S.A.U.

Marketing authorisation date:
1/12/2015

Manufacturing sites for batch release:
Laboratorios Maymo S.A.U.

Responsible authority:
Danish Medicines Agency

Authorisation number:
54739

Date of authorisation status change:
1/12/2015

Reference member state:
France

Procedure number:
FR/V/0282/001

Concerned member states:
Croatia Czechia Denmark Greece Hungary Ireland Italy Poland Portugal Romania 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

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

eu-puar-frv0282001-mr-rpe243-en.pdf