Aurofac Granular 250 mg/g Premix for medicated feedingstuff for pigs and chickens

Authorised

- Chlortetracycline hydrochloride

Product identification

Medicine name:
Aurofac Granular 250 mg/g Premix for medicated feedingstuff for pigs and chickens
Aurofac Granular 250 mg/g Premix for medicated feedingstuff for pigs and chickens

Active substance:
Chlortetracycline hydrochloride

Target species:
Chicken
Pig

Route of administration:
Oral use

Product details

Active substance and strength:
Chlortetracycline hydrochloride
250.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Premix for medicated feeding stuff

Withdrawal period by route of administration:

Oral use:

- **Chicken**
  - Eggs. 4 day
  - Meat and offal. 2 day
- **Pig**
  - Meat and offal. 10 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA03

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

Authorisation status:
Valid

Authorised in:
Ireland

Package description:
Polyethylene bags containing 2 kg
Polyethylene bags containing 3 kg
Polyethylene bags containing 4.8 kg
Polyethylene bags containing 6.4 kg
Polyethylene bags containing 8 kg
Polyethylene bags containing 9 kg
Polyethylene bags containing 12 kg
Polyethylene bags containing 16 kg
Polyethylene bags containing 20 kg
Polyethylene bags containing 25 kg
Polyethylene bags containing 3 kg. Cardboard cartons containing 8 x 3 kg.

Polyethylene bags containing 2 kg. Cardboard cartons containing 12 x 2 kg.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Phibro Animal Health (Poland) Sp. z o.o.

Marketing authorisation date:
9/12/2013

Manufacturing sites for batch release:
Zoetis Medolla Manufacturing S.r.l.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA25497/001/001

Date of authorisation status change:
9/12/2013

Reference member state:
Ireland

Procedure number:
IE/V/0207/002

Concerned member states:
Portugal Slovenia Spain

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

Documents

Summary of Product Characteristics