

# 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

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**Active substance:**

Chlortetracycline hydrochloride

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**Target species:**

Chicken

Pig

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Chlortetracycline hydrochloride

250.00 milligram(s) / 1.00 gram(s)

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**Pharmaceutical form:**

Premix for medicated feeding stuff

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**Withdrawal period by route of administration:****Oral use:**

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**Chicken**

- Eggs. 4 day
- Meat and offal. 2 day

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**Pig**

- Meat and offal. 10 day
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01AA03

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Portugal

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**Package description:**

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

Polyethylene bags containing 3 kg. Cardboard cartons containing 8 x 3 kg.

Polyethylene bags containing 25 kg

Polyethylene bags containing 20 kg

Polyethylene bags containing 16 kg

Polyethylene bags containing 12 kg

Polyethylene bags containing 9 kg

Polyethylene bags containing 8 kg

Polyethylene bags containing 6.4 kg

Polyethylene bags containing 4.8 kg

Polyethylene bags containing 3 kg

Polyethylene bags containing 2 kg

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Phibro Animal Health (Poland) Sp. z o.o.

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**Marketing authorisation date:**

31/08/2010

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**Manufacturing sites for batch release:**

Phibro Medolla Manufacturing S.r.l.

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**Responsible authority:**

Directorate General For Food And Veterinary

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**Authorisation number:**

275/01/10DFVPT

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**Date of authorisation status change:**

30/04/2026

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0207/002

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**Concerned member states:**

Portugal Slovenia Spain

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

Combined File of all Documents

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