# Amoxy Active, 697 mg/g oral powder for pigs and chickens

Authorised

- Amoxicillin trihydrate

## Product identification

**Medicine name:**
Amoxy Active, 697 mg/g oral powder for pigs and chickens
AMOXY ACTIVE 697 mg/g pulbere orala pentru porcine și pui de gaina

**Active substance:**
Amoxicillin trihydrate

**Target species:**
Chicken
Pig

**Route of administration:**
In drinking water use
In-feed use

## Product details

**Active substance and strength:**
Amoxicillin trihydrate
800.00 milligram(s) / 1.00 gram(s)

**Pharmaceutical form:**
Oral powder

**Withdrawal period by route of administration:**

**In drinking water use:**

- **Chicken**

  - Meat and offal. 1 day

- **Pig**

  - Meat and offal. 2 day

**In-feed use:**

- **Chicken**

  - Meat and offal. 1 day

- **Pig**

  - Meat and offal. 2 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01CA04

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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:**

Romania

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**Package description:**

500 g Securitainer: white polypropylene container , covered with a low-density polyethylene lid.
5 kg Bucket: white polypropylene bucket provided with a polypropylene lid.
250 g Securitainer: white polypropylene container , covered with a low-density polyethylene lid.
2,5 kg Bucket: white polypropylene bucket provided with a polypropylene lid.
100 g Securitainer: white polypropylene container , covered with a low-density polyethylene lid.
1 kg Securitainer: white polypropylene container , covered with a low-density polyethylene lid.
1 kg Bucket: white polypropylene bucket provided with a polypropylene lid.

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## 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:**
Dopharma Research B.V.

**Marketing authorisation date:**
1/06/2014

**Manufacturing sites for batch release:**
Dopharma B.V.

**Responsible authority:**
Institute For Control Of Biological Products And Veterinary Medicines

**Authorisation number:**
190088

**Date of authorisation status change:**
31/01/2024

**Reference member state:**
Netherlands

**Procedure number:**
NL/V/0179/001

**Concerned member states:**
Austria Belgium Bulgaria Croatia Czechia Denmark Estonia Finland France Germany Greece Hungary Ireland Italy Latvia Lithuania Norway Poland Portugal Romania Slovakia Sweden 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.

Combined File of all Documents

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