

# Amoxy Active CTD 697 mg/g Powderfor Use in Drinking Water for Chickens, Turkeys and Ducks

Authorised

- Amoxicillin trihydrate

## Product identification

**Medicine name:**

Amoxy Active CTD 697 mg/g Powderfor Use in Drinking Water for Chickens, Turkeys and Ducks

**Active substance:**

Amoxicillin trihydrate

**Target species:**

Chicken

Turkey

Duck

**Route of administration:**

In drinking water use

## Product details

**Active substance and strength:**

Amoxicillin trihydrate

800.00 milligram(s) / 1.00 gram(s)

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**Pharmaceutical form:**

Powder for use in drinking water

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**Withdrawal period by route of administration:**

**In drinking water use:**

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**Chicken**

- Meat and offal. 1 day

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**Turkey**

- Meat and offal. 5 day

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**Duck**

- Meat and offal. 9 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:**

France

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**Available in:**

France

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**Package description:**

250 g Securitainer: white cylindrical polypropylene container, covered with a low-density polyethylene closure.

2,5 kg Bucket: white polypropylene square container provided with a polypropylene closure.

500 g Securitainer: white cylindrical polypropylene container, covered with a low-density polyethylene closure.

5 kg Bucket: white polypropylene square container provided with a polypropylene closure.

100 g Securitainer: white cylindrical polypropylene container, covered with a low-density polyethylene closure.

1 kg Securitainer: white cylindrical polypropylene container, covered with a low-density polyethylene closure.

1 kg Bucket: white polypropylene square container provided with a polypropylene closure.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Dopharma Research B.V.

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**Marketing authorisation date:**

7/01/2019

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**Manufacturing sites for batch release:**

Dopharma B.V.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/3743369 3/2018

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**Date of authorisation status change:**

7/01/2019

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0308/001

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**Concerned member states:**

France Germany Hungary Italy Lithuania Poland Romania Spain  
United Kingdom (Northern Ireland)

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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

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.

### Combined File of all Documents