

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

Active substance:

Amoxicillin trihydrate

Target species:

Chicken

Pig

Route of administration:

In drinking water 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

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Pig

- Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

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.

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:

22/04/2014

Manufacturing sites for batch release:

Dopharma B.V.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/0924499 9/2014

Date of authorisation status change:

19/04/2019

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

Package Leaflet and Labelling

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