

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)

Pharmaceutical form:

Powder for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

-

Chicken

- Meat and offal. 1 day

-

Turkey

- Meat and offal. 5 day

-

Duck

- Meat and offal. 9 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Available in:

Italy

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.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Dopharma Research B.V.

Marketing authorisation date:

13/11/2019

Manufacturing sites for batch release:

Dopharma B.V.

Responsible authority:

Ministry Of Health

Authorisation number:

105209

Date of authorisation status change:

13/11/2019

Reference member state:

Netherlands

Procedure number:

NL/V/0308/001

Concerned member states:

France Germany Hungary Italy Lithuania Poland Romania Spain

United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)

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