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:
Lithuania

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:
25/02/2019

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/19/2522/001-007

Date of authorisation status change:
15/12/2025

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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