

Altidox 500 mg/g powder for use in drinking water for pigs, chickens and turkeys

Authorised

- Doxycycline hyclate

Product identification

Medicine name:

Altidox 500 mg/g powder for use in drinking water for pigs, chickens and turkeys

Active substance:

Doxycycline hyclate

Target species:

Turkey

Pig

Chicken

Route of administration:

In drinking water use

Product details

Active substance and strength:

Doxycycline hyclate

500.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water

Withdrawal period by route of administration:**In drinking water use:**

-

Turkey

- Meat and offal. 12 day

-

Pig

- Meat and offal. 4 day

-

Chicken

- Meat and offal. 5 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01AA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Available in:

Belgium

Package description:

2 kg bag with an outer layer of polyethylene terephthalate, middle layers of aluminium and polyamide and an inner layer of low density polyethylene (PET/ALU/PA/LDPE).

1 kg bag with an outer layer of polyethylene terephthalate, middle layers of aluminium and polyamide and an inner layer of low density polyethylene (PET/ALU/PA/LDPE).

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:

Dechra Regulatory B.V.

Marketing authorisation date:

17/11/2016

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V502737

Date of authorisation status change:

22/12/2021

Reference member state:

Netherlands

Procedure number:

NL/V/0277/001

Concerned member states:

Austria Belgium Croatia France Germany Hungary Ireland Italy Poland
Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 6/11/2025

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Package Leaflet

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

Package Leaflet and Labelling

Labelling

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