

# AMOXICILLIN GLOBAL VET

Authorised

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

- Amoxicillin trihydrate

## Product identification

### **Medicine name:**

AMOXICILLIN GLOBAL VET HEALTH 500 mg/g, powder for use in drinking water for chickens, turkeys, ducks and pigs

Amoksycylina Global Vet Health 500 mg/g Proszek do podania w wodzie do picia

### **Active substance:**

Amoxicillin trihydrate

### **Target species:**

Turkey

Chicken

Duck

Pig

### **Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Amoxicillin trihydrate  
500.00 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:****Oral use:**

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**Turkey**

- Meat and offal. 5 day

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**Chicken**

- Meat and offal. 1 day

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**Duck**

- Meat and offal. 9 day

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**Pig**

- Meat and offal. 2 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:**

Poland

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**Package description:**

The veterinary medicinal product is packed in thermo-sealed polyethylene / aluminium / polypropylene bags of 100 g.

The veterinary medicinal product is packed in thermo-sealed polyethylene / aluminium / polypropylene bags of 200 g.

The veterinary medicinal product is packed in thermo-sealed polyethylene / aluminium / polypropylene bags of 500 g.

The veterinary medicinal product is packed in thermo-sealed polyethylene / aluminium / polypropylene bags of 1 kg.

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**Additional information**

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Global Vet Health S.L.

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**Marketing authorisation date:**

1/07/2016

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**Manufacturing sites for batch release:**

S P Veterinaria S.A.

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

2544

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**Date of authorisation status change:**

1/07/2016

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**Reference member state:**

Ireland

**Procedure number:**

IE/V/0350/001

**Concerned member states:**

Bulgaria Cyprus France Greece Italy Malta Poland Portugal Romania Spain  
United Kingdom (Northern Ireland)

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

English (PDF)

Published on: 13/04/2025

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### Package Leaflet

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### Labelling

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