

File downloaded on 2026-04-15

**Source URL:** <https://medicines.health.europa.eu/veterinary/en/600000055171>

# CENVIMOX 500

Authorised

- Amoxicillin trihydrate

## Product identification

**Medicine name:**

CENVIMOX 500

---

**Active substance:**

Amoxicillin trihydrate

---

**Target species:**

Pig

Duck

Chicken (broiler)

Turkey (for meat production)

---

**Route of administration:**

In drinking water use

---

## Product details

**Active substance and strength:**

Amoxicillin trihydrate

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:**

- 

#### **Pig**

- Meat and offal. 6 day

- 

#### **Duck**

- Meat and offal. 7 day

- Eggs. no withdrawal period

Huevos: No autorizado en aves ponedoras para el consumo humano. No usar en un plazo de 4 semanas anteriores al comienzo del período de puesta ni durante la puesta

- 

#### **Chicken (broiler)**

- Meat and offal. 1 day

- Eggs. no withdrawal period

Huevos: No autorizado en aves ponedoras para el consumo humano. No usar en un plazo de 4 semanas anteriores al comienzo del período de puesta ni durante la puesta

- 

#### **Turkey (for meat production)**

- Meat and offal. 5 day

- Eggs. no withdrawal period

Huevos: No autorizado en aves ponedoras para el consumo humano. No usar en un plazo de 4 semanas anteriores al comienzo del período de puesta ni durante la puesta

---

## **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01CA04

---

## **Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Spain

---

**Package description:**

Available only in Spanish

Available only in Spanish

---

## 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:**

Cenavisa S.L.

---

**Marketing authorisation date:**

4/03/2008

---

**Manufacturing sites for batch release:**

Cenavisa S.L.

---

**Responsible authority:**

Spanish Agency For Medicines And Medical Devices

---

**Authorisation number:**

1850 ESP

---

**Date of authorisation status change:**

4/03/2008

---

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

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

### Package Leaflet

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

### Labelling

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