

# Clavoral 50/12.5 mg tablet for dogs and cats (strength 50 mg amoxicillin + 12.5 mg

Not authorised

- Clavulanic acid
- Amoxicillin trihydrate

## Product identification

### **Medicine name:**

Clavoral 50/12.5 mg tablet for dogs and cats (strength 50 mg amoxicillin + 12.5 mg CLAVUBACTIN 50/12,5 mg COMPRIMIDOS PARA GATOS Y PERROS

### **Active substance:**

Clavulanic acid

Amoxicillin trihydrate

### **Target species:**

Dog

Cat

### **Route of administration:**

Oral use

## Product details

### **Active substance and strength:**

Clavulanic acid  
12.50 milligram(s) / 1.00 Tablet  
Amoxicillin trihydrate  
50.00 milligram(s) / 1.00 Tablet

---

**Pharmaceutical form:**

Tablet

---

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

QJ01CR02

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Surrendered

---

**Authorised in:**

Spain

---

**Package description:**

Carton containing 5 aluminium/aluminium blister strips each strip with 4 tablets  
Carton containing 5 aluminium/aluminium blister strips each strip with 2 tablets.  
Carton containing 25 aluminium/aluminium blister strips each strip with 4 tablets.  
Carton containing 25 aluminium/aluminium blister strips each strip with 10 tablets  
Carton containing 10 aluminium/aluminium blister strips each strip with 10 tablets  
Carton containing 1 aluminium/aluminium blister strip with 10 tablets

---

## 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:**

Le Vet. B.V.

---

**Marketing authorisation date:**

30/11/2010

---

**Manufacturing sites for batch release:**

Lelypharma B.V.

---

**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

---

**Authorisation number:**

2221 ESP

---

**Date of authorisation status change:**

10/09/2025

---

**Reference member state:**

Netherlands

---

**Procedure number:**

NL/V/0149/001

---

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.