

# KESIUM 500 MG / 125 MG CHEWABLE TABLETS FOR DOGS

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

## Product identification

**Medicine name:**

KESIUM 500 MG / 125 MG CHEWABLE TABLETS FOR DOGS

Kesium 500 mg / 125 mg Kautabletten für Hunde

---

**Active substance:**

Amoxicillin trihydrate

Potassium clavulanate

---

**Target species:**

Dog

---

**Route of administration:**

Oral use

---

## Product details

**Active substance and strength:**

Amoxicillin trihydrate

574.00 milligram(s) / 1.00 Tablet

Potassium clavulanate

149.00 milligram(s) / 1.00 Tablet

---

**Pharmaceutical form:**

Chewable tablet

---

**Withdrawal period by route of administration:****Oral use:**

- Dog
- 

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

QJ01CR02

---

**Legal status of supply:**

Medicinal product on medical prescription for non-renewable delivery

---

**Authorisation status:**

Valid

---

**Authorised in:**

Austria

---

**Package description:**

Box of 80 blisters of 6 chewable breakable tablets

Box of 80 blisters of 6 chewable breakable tablets

Box of 80 blisters of 6 chewable breakable tablets

Box of 80 blisters of 6 chewable breakable tablets

Box of 80 blisters of 6 chewable breakable tablets

Box of 80 blisters of 6 chewable breakable tablets

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Ceva Sante Animale

---

**Marketing authorisation date:**

11/06/2013

---

**Manufacturing sites for batch release:**

Ceva Sante Animale

---

**Responsible authority:**

Austrian Agency For Health And Food Safety

---

**Authorisation number:**

8-01191

---

**Date of authorisation status change:**

24/04/2018

---

**Reference member state:**

France

---

**Procedure number:**

FR/V/0225/005

---

**Concerned member states:**

Austria Belgium Czechia Denmark Finland Germany Greece Hungary  
Ireland Italy Luxembourg Netherlands Poland Portugal Romania Spain  
Sweden 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

Package Leaflet

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

## Summary of Product Characteristics

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.

---

**Source URL:** <https://medicines.health.europa.eu/veterinary/600000029207>