

# 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+125 mg tyggetabletter

---

**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:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Denmark

---

**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:**

27/08/2013

---

**Manufacturing sites for batch release:**

Ceva Sante Animale

---

**Responsible authority:**

Danish Health And Medicines Authority

---

**Authorisation number:**

51087

---

**Date of authorisation status change:**

27/08/2013

---

**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)

---

**Source URL:** <https://medicines.health.europa.eu/veterinary/600000029138>