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# Clamaden 400 mg/100 mg chewable tablets for dogs

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

## Product identification

**Medicine name:**
Clamaden 400 mg/100 mg chewable tablets for dogs

**Active substance:**
Potassium clavulanate

Amoxicillin trihydrate

**Target species:**
Dog

**Route of administration:**
Oral use

## Product details

**Active substance and strength:**
Potassium clavulanate
119.13 milligram(s) / 1.00 Tablet

Amoxicillin trihydrate

459.22 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**
Chewable tablet

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QJ01CR02

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Latvia

**Available in:**
Latvia

**Package description:**
Blister formed of an aluminium foil which consists of aluminum layer coated with OPA (Oriented Polyamide) film on one side and PE with desiccant on the other side and an aluminium sealing foil which consists of aluminium layer and PE coating.Blister contains 6 tablets. Carton contains 12 tablets
Blister formed of an aluminium foil which consists of aluminum layer coated with OPA (Oriented Polyamide) film on one side and PE with desiccant on the other side and an aluminium sealing foil which consists of aluminium layer and PE coating.Blister contains 6 tablets. Carton contains 60 tablets
Blister formed of an aluminium foil which consists of aluminum layer coated with OPA (Oriented Polyamide) film on one side and PE with desiccant on the other side and an aluminium sealing foil which consists of aluminium layer and PE coating.Blister contains 6 tablets. Carton contains 300 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:**
KRKA tovarna zdravil d.d. Novo mesto

**Marketing authorisation date:**
23/02/2022

**Manufacturing sites for batch release:**
KRKA tovarna zdravil d.d. Novo mesto

**Responsible authority:**
Food And Veterinary Service

**Authorisation number:**
V/MRP/22/0008

**Date of authorisation status change:**
23/02/2022

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0652/003

**Concerned member states:**
Belgium Croatia Czechia Estonia France Germany Hungary Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

# Documents

Summary of Product Characteristics

Combined File of all Documents

English (PDF)
Published on: 20/09/2026
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