

KESIUM 40 MG / 10 MG CHEWABLE TABLETS FOR CATS AND DOGS

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:

KESIUM 40 MG / 10 MG CHEWABLE TABLETS FOR CATS AND DOGS

Active substance:

Amoxicillin trihydrate

Potassium clavulanate

Target species:

Dog

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Amoxicillin trihydrate

45.90 milligram(s) / 1.00 Tablet

Potassium clavulanate

11.90 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:

Greece

Package description:

Box of 1 blister (PA-AL-PVC - aluminium heat sealed) of 10 tablets

Box of 24 blisters (PA-AL-PVC - aluminium heat sealed) of 10 tablets

Box of 10 blisters (PA-AL-PVC - aluminium heat sealed) of 10 tablets

Box of 8 blisters (PA-AL-PVC - aluminium heat sealed) of 10 tablets

Box of 6 blisters (PA-AL-PVC - aluminium heat sealed) of 10 tablets

Box of 4 blisters (PA-AL-PVC - aluminium heat sealed) of 10 tablets

Box of 2 blisters (PA-AL-PVC - aluminium heat sealed) of 10 tablets

Box of 48 blisters (PA-AL-PVC - aluminium heat sealed) of 10 tablets

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:

Ceva Hellas LLC

Marketing authorisation date:

10/06/2012

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

National Organization For Medicines

Authorisation number:

2028/11-01-2017/K-0191101

Date of authorisation status change:

10/01/2017

Reference member state:

France

Procedure number:

FR/V/0225/001

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