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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

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CR02

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

Authorisation status:
Valid

Authorised in:
Greece

Available in:
Greece

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Box of 80 blisters (PA-AL-PVC – aluminium heat sealed) of 6 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 Hellas LLC

Marketing authorisation date:

11/03/2014

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
National Organization For Medicines

Authorisation number:
95915/17/04-10-2018/K-0191105

Date of authorisation status change:
3/10/2018

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

Documents

Combined File of all Documents