

KESIUM 400 MG / 100 MG CHEWABLE TABLETS FOR DOGS

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:

KESIUM 400 MG / 100 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

459.20 milligram(s) / 1.00 Tablet

Potassium clavulanate

119.10 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:

Italy

Package description:

Available only in [French](#)

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Box of 80 blisters (PA-AL-PVC – aluminium heat sealed) of 6 tablets

Box of 3 blisters (PA-AL-PVC – aluminium heat sealed) of 4 tablets

Box of 6 blisters (PA-AL-PVC – aluminium heat sealed) of 4 tablets

Box of 9 blisters (PA-AL-PVC – aluminium heat sealed) of 4 tablets

Box of 12 blisters (PA-AL-PVC – aluminium heat sealed) of 4 tablets

Box of 15 blisters (PA-AL-PVC – aluminium heat sealed) of 4 tablets

Box of 18 blisters (PA-AL-PVC – aluminium heat sealed) of 4 tablets

Box of 21 blisters (PA-AL-PVC – aluminium heat sealed) of 4 tablets

Box of 24 blisters (PA-AL-PVC – aluminium heat sealed) of 4 tablets

Box of 60 blisters (PA-AL-PVC – aluminium heat sealed) of 4 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 Salute Animale S.p.A.

Marketing authorisation date:

9/03/2012

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

Ministry Of Health

Authorisation number:

104319

Date of authorisation status change:

9/03/2012

Reference member state:

France

Procedure number:

FR/V/0225/004

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

English (PDF)

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