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:
Finland

Available in:
Finland

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
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
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 Sante Animale

Marketing authorisation date:
11/12/2011

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Finnish Medicines Agency

Authorisation number:
28974

Date of authorisation status change:
11/12/2011

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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents