KESIUM 50 MG / 12,5 MG CHEWABLE TABLETS FOR CATS AND DOGS

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:
KESIUM 50 MG / 12,5 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

57.40 milligram(s) / 1.00 Tablet

Potassium clavulanate
14.90 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:
Poland

Available in:
Poland

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:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Animal Health Polska Sp. z o.o.

Marketing authorisation date:
13/04/2012

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2174

Date of authorisation status change:
13/04/2012

Reference member state:
France

Procedure number:
FR/V/0225/002

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.

Labelling

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

Combined File of all Documents