KESIUM 200 MG / 50 MG CHEWABLE TABLETS FOR DOGS

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:
KESIUM 200 MG / 50 MG CHEWABLE TABLETS FOR DOGS
Kesium 200 mg/50 mg Tuggtablett

Active substance:
Amoxicillin trihydrate
Potassium clavulanate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Amoxicillin trihydrate
229.60 milligram(s) / 1.00 Tablet

Potassium clavulanate
59.50 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:
Sweden

Available in:
Sweden

Package description:
Box of 1 blister (PA-AL-PVC – aluminium heat sealed) of 8 tablets
Box of 30 blisters (PA-AL-PVC – aluminium heat sealed) of 8 tablets
Box of 12 blisters (PA-AL-PVC – aluminium heat sealed) of 8 tablets
Box of 10 blisters (PA-AL-PVC – aluminium heat sealed) of 8 tablets
Box of 8 blisters (PA-AL-PVC – aluminium heat sealed) of 8 tablets
Box of 6 blisters (PA-AL-PVC – aluminium heat sealed) of 8 tablets
Box of 4 blisters (PA-AL-PVC – aluminium heat sealed) of 8 tablets
Box of 2 blisters (PA-AL-PVC – aluminium heat sealed) of 8 tablets
Box of 60 blisters (PA-AL-PVC – aluminium heat sealed) of 8 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:
30/09/2011

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
44787

Date of authorisation status change:
30/09/2011

Reference member state:
France

Procedure number:
FR/V/0225/003

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

Package Leaflet

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Summary of Product Characteristics

English (PDF)
Published on: 24/09/2024
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