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
Kesium 500 mg/125 mg rágótabletta kutyák részére A.U.V.

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

Withdrawal period by route of administration:

Oral use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CR02

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

Authorisation status:
Valid

Authorised in:
Hungary

Package description:
Box of 80 blisters of 6 chewable breakable tablets
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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-Phylaxia Zrt.

Marketing authorisation date:

25/09/2013

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
3417/X/13 NÉBIH ÁTI

Date of authorisation status change:
25/09/2013

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

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