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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:
Germany

Available in:
Germany

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
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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 Tiergesundheit GmbH

Marketing authorisation date:
4/08/2011

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401476.03.00

Date of authorisation status change:
28/07/2016

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)

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Documents

Combined File of all Documents

English (PDF)
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