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Clamaden 40 mg/10 mg chewable tablets for cats and dogs

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
Clamaden 40 mg/10 mg chewable tablets for cats and dogs

Active substance:
Potassium clavulanate
Amoxicillin trihydrate

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Potassium clavulanate
11.91 milligram(s) / 1.00 Tablet

Amoxicillin trihydrate
45.92 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:
Lithuania

Available in:
Lithuania

Package description:
Blister formed of an aluminium foil which consists of aluminum layer coated with OPA (Oriented Polyamide) film on one side and PE with desiccant on the other side and an aluminium sealing foil which consists of aluminium layer and PE coating.Blister contains 10 tablets. Carton contains 500 tablets.
Blister formed of an aluminium foil which consists of aluminum layer coated with OPA (Oriented Polyamide) film on one side and PE with desiccant on the other side and an aluminium sealing foil which consists of aluminium layer and PE coating.Blister contains 10 tablets. Carton contains 100 tablets.
Blister formed of an aluminium foil which consists of aluminum layer coated with OPA (Oriented Polyamide) film on one side and PE with desiccant on the other side and an aluminium sealing foil which consists of aluminium layer and PE coating.Blister contains 10 tablets. Carton contains 20 tablets.
Blister formed of an aluminium foil which consists of aluminum layer coated with OPA (Oriented Polyamide) film on one side and PE with desiccant on the other side and an aluminium sealing foil which consists of aluminium layer and PE coating.Blister contains 10 tablets. Carton contains 10 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:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
6/03/2022

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/22/2700/001-004

Date of authorisation status change:
26/10/2026

Reference member state:
Ireland

Procedure number:
IE/V/0652/001

Concerned member states:
Belgium Croatia Czechia Estonia France Germany Hungary Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

English (PDF)
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Summary of Product Characteristics