Clavudale 40 mg/10 mg tablets for cats and dogs

Authorised

- Amoxicillin
- Potassium clavulanate

Product identification

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

Active substance:
Amoxicillin
Potassium clavulanate

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

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

Potassium clavulanate
11.91 milligram(s) / 1.00 Tablet

Pharmaceutical form:
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

Package description:
Blister packs consisting of orientated Polyamide/Aluminium/Polyvinyl chloride film, heat sealed with aluminium foil(25micrometer) in strips of 6 tablets. Cartons containing 120 tablets.
Blister packs consisting of orientated Polyamide/Aluminium/Polyvinyl chloride film, heat sealed with aluminium foil(25micrometer) in strips of 6 tablets. Cartons containing 24 tablets.
Blister packs consisting of orientated Polyamide/Aluminium/Polyvinyl chloride film, heat sealed with aluminium foil(25micrometer) in strips of 6 tablets. Cartons containing 12 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:
Dechra Regulatory B.V.

Marketing authorisation date:
28/07/2020

Manufacturing sites for batch release:
Genera d.d.
Laboratorio Reig Jofre S.A.

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
60199

Date of authorisation status change:
28/07/2020

Reference member state:
Ireland

Procedure number:
IE/V/0504/001

Concerned member states:
Austria Belgium Croatia Czechia Denmark Finland France Germany Greece Hungary Iceland Italy Luxembourg Netherlands Norway Poland Portugal Slovakia Slovenia 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)
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