Clavudale 40 mg/10 mg tablets for cats and dogs

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:
Clavudale 40 mg/10 mg tablets for cats and dogs
Clavudale 40 mg/10 mg tabletta kutyák és macskák részére A.U.V

Active substance:
Amoxicillin trihydrate

Potassium clavulanate

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

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

Potassium clavulanate

11.91 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Withdrawal period by route of administration:
Oral use:

- **Dog**
- **Cat**

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:
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.
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 120 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:
19/10/2011

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

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
3014/X/11 MgSzH ÁTI

Date of authorisation status change:
19/10/2011

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

Summary of Product Characteristics
