

Clavudale 200 mg/50 mg tablets for dogs

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:

Clavudale 200 mg/50 mg 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

229.61 milligram(s) / 1.00 Tablet

Potassium clavulanate

59.56 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:

Finland

Available in:

Finland

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:

17/09/2020

Manufacturing sites for batch release:

Genera d.d.

Laboratorio Reig Jofre S.A.

Responsible authority:

Finnish Medicines Agency

Authorisation number:

37794

Date of authorisation status change:

17/09/2020

Reference member state:

Ireland

Procedure number:

IE/V/0504/002

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

English (PDF)

Published on: 3/05/2024

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Package Leaflet

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