Source URL: https://medicines.health.europa.eu/veterinary/en/60000050918

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 Spectrabactin Vet 200 mg/50 mg Tablett

Active substance:

Amoxicillin trihydrate

Potassium clavulanate

Target species:

Dog

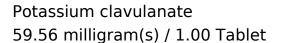
Route of administration:

Oral use

Product details

Active substance and strength:

Amoxicillin trihydrate 229.61 milligram(s) / 1.00 Tablet



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 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: 28/07/2020
Manufacturing sites for batch release: Genera d.d.
Laboratorio Reig Jofre S.A.
Responsible authority: Swedish Medical Products Agency
Authorisation number: 60200
Date of authorisation status change: 28/07/2020
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
Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

English (PDF)

Published on: 3/05/2024

Download

Labelling

This document does not exist in this language (English). You can find it in another language below.