

Clavucill 200 mg + 50 mg 200 mg - 50 mg Tablet

Authorised

- Potassium clavulanate, diluted (potassium clavulanate - cellulose, microcrystalline (1:1))
- Amoxicillin trihydrate

Product identification

Medicine name:

Clavucill 200 mg + 50 mg 200 mg - 50 mg Tablet

Active substance:

Potassium clavulanate, diluted (potassium clavulanate - cellulose, microcrystalline (1:1))

Amoxicillin trihydrate

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Potassium clavulanate, diluted (potassium clavulanate - cellulose, microcrystalline (1:1))

125.10 milligram(s) / 1.00 Tablet

Amoxicillin trihydrate

229.55 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:

Denmark

Package description:

Clavucill 200 mg + 50 mg 200 mg - 50 mg tabl. 250

Clavucill 200 mg + 50 mg 200 mg - 50 mg tabl. 100

Clavucill 200 mg + 50 mg 200 mg - 50 mg tabl. 10

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:

V.M.D.

Marketing authorisation date:

20/10/2011

Manufacturing sites for batch release:

V.M.D.

Responsible authority:

Danish Medicines Agency

Authorisation number:

48710

Date of authorisation status change:

20/10/2011

Reference member state:

Belgium

Procedure number:

BE/V/0024/002

Concerned member states:

Denmark France Germany Netherlands Poland Portugal Romania Spain
Sweden

Generic of:

600000085703

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)

Published on: 13/03/2026

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Combined File of all Documents

Package Leaflet

English (PDF)

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