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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

Available in:
Denmark

Package description:
Clavucill 200 mg + 50 mg 200 mg - 50 mg tabl. 10
Clavucill 200 mg + 50 mg 200 mg - 50 mg tabl. 100
Clavucill 200 mg + 50 mg 200 mg - 50 mg tabl. 250

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: 9/07/2026

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Combined File of all Documents

Package Leaflet

Labelling