

Clavoral 250/62.5 mg tablet for dogs and cats (strength 250 mg amoxicillin + 62.5 mg)

Not authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:

Clavoral 250/62.5 mg tablet for dogs and cats (strength 250 mg amoxicillin + 62.5 mg)

Active substance:

Potassium clavulanate
Amoxicillin trihydrate

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Potassium clavulanate

74.45 milligram(s) / 1.00 Tablet

Amoxicillin trihydrate

287.00 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:

Surrendered

Authorised in:

Netherlands

Package description:

Carton containing 5 aluminium/aluminium blister strips each strip with 4 tablets corresponding to 20 tablets per carton.

Carton containing 5 aluminium/aluminium blister strips each strip with 2 tablets corresponding to 10 tablets per carton.

Carton containing 25 aluminium/aluminium blister strips each strip with 4 tablets corresponding to 100 tablets per carton

Carton containing 25 aluminium/aluminium blister strips each strip with 10 tablets corresponding to 250 tablets per carton

Carton containing 10 aluminium/aluminium blister strips each strip with 10 tablets corresponding to 100 tablets per carton

Carton containing 1 aluminium/aluminium blister strip with 10 tablets corresponding to 10 tablets per carton.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Ast Farma B.V.

Marketing authorisation date:

30/07/2003

Manufacturing sites for batch release:

Lelypharma B.V.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 10086

Date of authorisation status change:

18/09/2025

Reference member state:

Netherlands

Procedure number:

NL/V/0149/002

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

Documents

Combined File of all Documents

English (PDF)

Published on: 27/10/2025

Download