Clavoral 500/125 mg tablet for dogs (strength 500 mg amoxicillin + 125 mg

Not authorised

- Amoxicillin trihydrate
- Clavulanic acid

Product identification

Medicine name:
Clavoral 500/125 mg tablet for dogs (strength 500 mg amoxicillin + 125 mg
CLAVUBACTIN 500/125 mg COMPRIMIDOS PARA PERROS

Active substance:
Amoxicillin trihydrate

Clavulanic acid

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Amoxicillin trihydrate

500.00 milligram(s) / 1.00 Tablet

Clavulanic acid
125.00 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:
Spain

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:
Le Vet. B.V.

Marketing authorisation date:
30/11/2010

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
2223 ESP

Date of authorisation status change:
10/09/2025

Reference member state:
Netherlands

Procedure number:
NL/V/0149/003

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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