

Amoxicillin 500 mg tablets for dogs

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:

Amoxicillin 500 mg tablets for dogs

Active substance:

Amoxicillin trihydrate

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Amoxicillin trihydrate

575.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Available in:

Portugal

Package description:

Cardboard box containing 10 separate cardboard boxes, each containing 1 blister of 10 tablets

Cardboard box of 2 Aluminium - PVC/PE/PVDC blisters of 10 tablets.

Cardboard box of 25 Aluminium - PVC/PE/PVDC blisters of 10 tablets

Cardboard box of 3 Aluminium - PVC/PE/PVDC blisters of 10 tablets.

Cardboard box of 4 Aluminium - PVC/PE/PVDC blisters of 10 tablets.

Cardboard box of 5 Aluminium - PVC/PE/PVDC blisters of 10 tablets.

Cardboard box of 50 Aluminium - PVC/PE/PVDC blisters of 10 tablets

Cardboard box of 6 Aluminium - PVC/PE/PVDC blisters of 10 tablets.

Cardboard box of 7 Aluminium - PVC/PE/PVDC blisters of 10 tablets.

Cardboard box of 8 Aluminium - PVC/PE/PVDC blisters of 10 tablets.

Cardboard box of 9 Aluminium - PVC/PE/PVDC blisters of 10 tablets.

Cardboard box of 1 Aluminium - PVC/PE/PVDC blister of 10 tablets.

Cardboard box of 10 Aluminium - PVC/PE/PVDC blisters of 10 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:

Le Vet. Beheer B.V.

Marketing authorisation date:

14/10/2014

Manufacturing sites for batch release:

Lelypharma B.V.

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

848/03/14DFVPT

Date of authorisation status change:

6/05/2026

Reference member state:

Netherlands

Procedure number:

NL/V/0186/003

Concerned member states:

Austria Belgium Cyprus Czechia Denmark Estonia Finland France Germany
Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway
Poland Portugal Romania Slovakia Spain United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

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