

# Amoxicillin 250 mg tablets for dogs

Authorised

- Amoxicillin trihydrate

## Product identification

**Medicine name:**

Amoxicillin 250 mg tablets for dogs

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**Active substance:**

Amoxicillin trihydrate

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Amoxicillin trihydrate

287.50 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01CA04

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Finland

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**Package description:**

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

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

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

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

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

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

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

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

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

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

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

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

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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Le Vet. Beheer B.V.

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**Marketing authorisation date:**

1/04/2015

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**Manufacturing sites for batch release:**

Lelypharma B.V.

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**Responsible authority:**

Finnish Medicines Agency

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**Authorisation number:**

31904

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**Date of authorisation status change:**

1/04/2015

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0186/002

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**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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[www.adrreports.eu/vet](http://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.