

# Amoxibactin 250 mg tablets for dogs

Authorised

- Amoxicillin trihydrate

## Product identification

**Medicine name:**

Amoxibactin 250 mg tablets for dogs

Amoxibactin 250 mg compresse per cani

**Active substance:**

Amoxicillin trihydrate

**Target species:**

Dog

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Amoxicillin trihydrate

287.50 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**

Tablet

**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:**

Italy

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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:**

12/02/2016

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**Manufacturing sites for batch release:**

Lelypharma B.V.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

104712

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**Date of authorisation status change:**

24/09/2019

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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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## Documents

Combined File of all Documents

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