

# Cylanic 250 mg + 62.5 mg tablets for dogs and cats

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

## Product identification

### Medicine name:

Cylanic 250 mg + 62.5 mg tablets for dogs and cats

Cylanic 250 mg + 62,5 mg Tabletká

### Active substance:

Potassium clavulanate

Amoxicillin trihydrate

### Target species:

Dog

Cat

### Route of administration:

Oral use

## Product details

### Active substance and strength:

Potassium clavulanate

74.45 milligram(s) / 1.00 Tablet

Amoxicillin trihydrate  
287.01 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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

QJ01CR02

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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:**

Poland

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**Package description:**

oPA/Alu/PVC - PVC/Alu heat sealed blister containing 10 tablets each. Package sizes: Cardboard box of 250 tablets.

oPA/Alu/PVC - PVC/Alu heat sealed blister containing 10 tablets each. Package sizes: Cardboard box of 100 tablets.

oPA/Alu/PVC - PVC/Alu heat sealed blister containing 10 tablets each. Package sizes: Cardboard box of 50 tablets.

oPA/Alu/PVC - PVC/Alu heat sealed blister containing 10 tablets each. Package sizes: Cardboard box of 30 tablets.

oPA/Alu/PVC - PVC/Alu heat sealed blister containing 10 tablets each. Package sizes: Cardboard box of 10 tablets.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Industrial Veterinaria S.A.

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**Marketing authorisation date:**

20/07/2021

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**Manufacturing sites for batch release:**

Industrial Veterinaria S.A.  
aniMedica Herstellungs GmbH  
Lelypharma B.V.  
aniMedica GmbH

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

3117

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**Date of authorisation status change:**

20/07/2021

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0582/002

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia Germany Greece  
Hungary Italy Latvia Lithuania Netherlands Poland Portugal Romania  
Slovakia Slovenia Spain

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Summary of Product Characteristics

English (PDF)

Published on: 11/02/2022

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### Package Leaflet

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### Labelling

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