

Cylanic 50 mg + 12.5 mg tablets for dogs and cats

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:

Cylanic 50 mg + 12.5 mg tablets for dogs and cats

Cylanic 50 mg + 12,5 mg Tabletká

Active substance:

Amoxicillin trihydrate

Potassium clavulanate

Target species:

Dog

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Amoxicillin trihydrate

57.40 milligram(s) / 1.00 Tablet

Potassium clavulanate
14.89 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:

Valid

Authorised in:

Poland

Available in:

Poland

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.

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:

Industrial Veterinaria S.A.

Marketing authorisation date:

20/07/2021

Manufacturing sites for batch release:

Lelypharma B.V.

aniMedica GmbH

Industrial Veterinaria S.A.

aniMedica Herstellungs GmbH

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

3116

Date of authorisation status change:

20/07/2021

Reference member state:

Ireland

Procedure number:

IE/V/0582/001

Concerned member states:

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

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 25/01/2026

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Package Leaflet

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Labelling

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