

# Clindabactin 440 mg Chewable Tablets for Dogs

Authorised

- Clindamycin hydrochloride

## Product identification

**Medicine name:**

Clindabactin 440 mg Chewable Tablets for Dogs

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**Active substance:**

Clindamycin hydrochloride

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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:**

Clindamycin hydrochloride  
440.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Chewable tablet

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

QJ01FF01

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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:**

Romania

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**Available in:**

Romania

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**Package description:**

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box containing 10 separate cardboard boxes, each containing 1 blister of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 8 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 6 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 4 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 3 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 25 blisters of 10 tablets.

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

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 9 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 7 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 5 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 2 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 10 blisters 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:**

Dechra Regulatory B.V.

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**Marketing authorisation date:**

4/06/2019

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**Manufacturing sites for batch release:**

Lelypharma B.V.

Genera d.d.

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**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

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**Authorisation number:**

240071

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**Date of authorisation status change:**

16/06/2025

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0317/003

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Hungary Iceland Ireland Italy Latvia Lithuania

Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain  
Sweden United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
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## Documents

Combined File of all Documents

English (PDF)

Published on: 24/03/2025

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