

# Kefavet® vet 250 mg, film-coated tablet

Authorised

- Cefalexin monohydrate

## Product identification

**Medicine name:**

Kefavet® vet 250 mg, film-coated tablet  
KEFAVET vet. 250 mg comprimate filmate

**Active substance:**

Cefalexin monohydrate

**Target species:**

Dog

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Cefalexin monohydrate  
263.00 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**

Film-coated tablet

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

QJ01DB01

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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:**

Available only in [Swedish](#)

Available only in [Swedish](#)

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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:**

Orion Corporation

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**Marketing authorisation date:**

17/08/2010

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**Manufacturing sites for batch release:**

Orion Oyj / Orion Corporation / Orion Pharma

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**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

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**Authorisation number:**

110050

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**Date of authorisation status change:**

7/05/2025

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**Reference member state:**

Sweden

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**Procedure number:**

SE/V/0114/001

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**Concerned member states:**

Belgium Czechia Denmark Estonia Hungary Iceland Latvia Lithuania  
Luxembourg Netherlands Poland Romania Slovakia Slovenia

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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

Combined File of all Documents

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

Package Leaflet

Summary of Product Characteristics

Labelling