# 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 plėvele dengtos tabletės

**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

**Withdrawal period by route of administration:**

**Oral use:**

- **Dog**

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

QJ01DB01

**Legal status of supply:**

This information is not available for this product.

**Authorisation status:**

Valid

**Authorised in:**

Lithuania

**Available in:**

Lithuania

**Package description:**

Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish

## 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:**

Orion Corporation

**Marketing authorisation date:**

30/06/2010

**Manufacturing sites for batch release:**
Orion Corporation

**Responsible authority:**
State Food And Veterinary Service

**Authorisation number:**
LT/2/10/1943/001-005

**Date of authorisation status change:**
26/01/2011

**Reference member state:**
Sweden

**Procedure number:**
SE/V/0114/001

**Concerned member states:**
Belgium Czechia Denmark Estonia Hungary Iceland Latvia Lithuania Luxembourg Netherlands Poland Romania Slovakia Slovenia

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

## Documents

Summary of Product Characteristics

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

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