

# SEMELCEF 1000 mg tablets for dogs

Authorised

- Cefadroxil monohydrate

## Product identification

**Medicine name:**

SEMELCEF 1000 mg tablets for dogs

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**Active substance:**

Cefadroxil monohydrate

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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:**

Cefadroxil monohydrate

1050.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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

QJ01DB05

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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:**

Slovakia

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**Available in:**

Slovakia

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**Package description:**

box containing 10 blisters of 6 tablets (60 tablets)

box containing 1 blister of 6 tablets

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Fatro S.p.A.

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**Marketing authorisation date:**

18/12/2019

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**Manufacturing sites for batch release:**

Fatro S.p.A.

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/057/DC/19-S

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**Date of authorisation status change:**

18/12/2019

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0304/002

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**Concerned member states:**

Austria Belgium Croatia Cyprus Czechia Estonia Greece Hungary Ireland  
Poland Portugal 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.

Labelling

Summary of Product Characteristics

Package Leaflet

eu-PUAR-esv0304002-dcp-semelcef-1000-mg-tablets-for-dogs-en.pdf