

SEMELCEF 1000 mg tablets for dogs

Authorised

- Cefadroxil monohydrate

Product identification

Medicine name:

SEMELCEF 1000 mg tablets for dogs

Active substance:

Cefadroxil monohydrate

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Cefadroxil monohydrate
1050.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01DB05

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Package description:

box containing 10 blisters of 6 tablets (60 tablets)

box containing 1 blister of 6 tablets

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Fatro S.p.A.

Marketing authorisation date:

13/03/2020

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

2960

Date of authorisation status change:

13/03/2020

Reference member state:

Spain

Procedure number:

ES/V/0304/002

Concerned member states:

Austria Belgium Croatia Cyprus Czechia Estonia Greece Hungary Ireland
Poland Portugal Slovakia Slovenia

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

Documents

Package Leaflet

English (PDF)

Published on: 29/01/2026

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Summary of Product Characteristics

English (PDF)

Published on: 29/01/2026

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Labelling

English (PDF)

Published on: 22/12/2023

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