

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:

Croatia

Available in:

Croatia

Package description:

box containing 1 blister of 6 tablets

box containing 10 blisters of 6 tablets (60 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:

28/06/2019

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/19-01/415

Date of authorisation status change:

13/02/2026

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

Summary of Product Characteristics

English (PDF)

Published on: 29/01/2026

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Labelling

Package Leaflet

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