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:
Estonia

Available in:
Estonia

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:
1/07/2019

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
State Agency Of Medicines

Authorisation number:

2185

Date of authorisation status change:
1/07/2019

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

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