

File downloaded on 2026-05-11

Source URL: <https://medicines.health.europa.eu/veterinary/en/600000023260>

Kefavet® vet 250 mg, film-coated tablet

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:

Kefavet® vet 250 mg, film-coated tablet

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01DB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Sweden

Available in:

Sweden

Package description:

Blister, 28 tablets

Blister, 70 tablets

Blister, 140 tablets

Blister, 14 tablets

Blister, 20 tablets

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:

22/09/2000

Manufacturing sites for batch release:

Orion Corporation

Responsible authority:

Swedish Medical Products Agency

Authorisation number:

13808

Date of authorisation status change:

22/09/2000

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

Package Leaflet

English (PDF)

Published on: 17/04/2025

[Download](#)

Summary of Product Characteristics

English (PDF)

Published on: 17/04/2025

[Download](#)

Labelling

English (PDF)

Published on: 17/04/2025

[Download](#)