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Kefavet® vet 500 mg, film-coated tablet

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:
Kefavet® vet 500 mg, film-coated tablet

Active substance:
Cefalexin monohydrate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Cefalexin monohydrate
526.00 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, 140 tablets
Blister, 70 tablets
Blister, 14 tablets
Blister, 30 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:
13810

Date of authorisation status change:
22/09/2000

Reference member state:
Sweden

Procedure number:
SE/V/0114/002

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
Updated on: 14/03/2026
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Summary of Product Characteristics

English (PDF)
Published on: 17/04/2025
Updated on: 14/03/2026
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Labelling

English (PDF)
Published on: 17/04/2025
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