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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:

Czechia

Available in:

Czechia

Package description:

Available only in [Swedish](#)

Available only in [Swedish](#)

Available only in [Swedish](#)

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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:

17/03/2008

Manufacturing sites for batch release:

Orion Corporation

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/015/08-C

Date of authorisation status change:

17/03/2008

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

Summary of Product Characteristics

English (PDF)

Published on: 17/04/2025

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Package Leaflet

English (PDF)

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Labelling

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