

Kefavet® vet. 500 mg, film-coated tablet

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:

Kefavet® vet. 500 mg, film-coated tablet
Kefavet vet. 500 mg apvalkotās tabletes suņiem

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

Withdrawal period by route of administration:

Oral use:

- Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01DB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

Available in:

Latvia

Package description:

Available only in [Latvian](#)

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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:

3/08/2010

Manufacturing sites for batch release:

Orion Corporation

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/MRP/10/0023

Date of authorisation status change:

3/08/2010

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

Source URL: <https://medicines.health.europa.eu/veterinary/600000023298>