Kefavet® vet. 250 mg, film-coated tablet

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:
Kefavet® vet. 250 mg, film-coated tablet
Kefavet Vet. 250 mg filmovertrukne tabletter

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

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:

Denmark

Available in:

Denmark

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:

11/11/2013

Manufacturing sites for batch release:
Orion Corporation

Responsible authority:
Danish Health And Medicines Authority

Authorisation number:
52199

Date of authorisation status change:
11/11/2013

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

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