Rilexine, 300 mg tabletid koertele

Authorised

- Cefalexin

Product identification

Medicine name:
Rilexine, 300 mg tabletid koertele

Active substance:
Cefalexin

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Cefalexin
300.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
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:
Estonia

Available in:
Estonia

Package description:
Available only in Estonian
Available only in Estonian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Virbac

Marketing authorisation date:
14/12/2006

Manufacturing sites for batch release:
Virbac S.A.

Responsible authority:
State Agency Of Medicines

Authorisation number:
1444

Date of authorisation status change:
14/12/2006

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.