

RILEXINE 300 mg, tabletès šunims

Authorised

- Cefalexin

Product identification

Medicine name:

RILEXINE 300 mg, tabletès šunims

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01DB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Lithuania

Available in:

Lithuania

Package description:

Available only in Lithuanian

Available only in Lithuanian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Virbac

Marketing authorisation date:

26/11/2006

Manufacturing sites for batch release:

Virbac S.A.

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/06/1716/001-002

Date of authorisation status change:

26/12/2011

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

Documents

RV1716.pdf