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Rilexine 600 600 mg Tablet

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:
Rilexine 600 600 mg Tablet

Active substance:
Cefalexin monohydrate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Cefalexin monohydrate
631.11 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:
Belgium

Available in:
Belgium

Package description:
Rilexine 600 600 mg Tablet Box with 2 blisters of 7 tablets
Rilexine 600 600 mg Tablet Box with 3 blisters of 7 tablets
Rilexine 600 600 mg Tablet Box with 4 blisters of 7 tablets
Rilexine 600 600 mg Tablet Box with 20 blisters of 7 tablets
Rilexine 600 600 mg Tablet Box with 30 blisters of 7 tablets

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:
19/07/2004

Manufacturing sites for batch release:
Virbac

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V265614

Date of authorisation status change:
12/02/2018

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

Documents

Package Leaflet

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Summary of Product Characteristics

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

Labelling

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