Rilexine 75 mg, tablete, za pse i mačke

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:
Rilexine 75 mg, tablete, za pse i mačke

Active substance:
Cefalexin monohydrate

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Cefalexin monohydrate
75.00 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:
Croatia

Package description:
Available only in Croatian
Available only in Croatian
Available only in Croatian

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:
18/06/2018

Manufacturing sites for batch release:
Virbac

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/12-01/356

Date of authorisation status change:
29/07/2025

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

Documents

Summary of Product Characteristics

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