

CEFASEPTIN 75 MG TABLETS FOR DOGS AND CATS

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:

CEFASEPTIN P 75MG ΔΙΣΚΙΟ
CEFASEPTIN 75 MG TABLETS FOR DOGS AND CATS

Active substance:

Cefalexin monohydrate

Target species:

Dog
Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Cefalexin monohydrate
78.90 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Withdrawal period by route of administration:**Oral use:**

-

Dog

-

Cat

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01DB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Package description:Available only in [French](#)Available only in [French](#)Available only in [French](#)

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:

Vetoquinol S.A.

Marketing authorisation date:

6/03/2016

Manufacturing sites for batch release:

Vetoquinol S.A.

Responsible authority:

National Organization For Medicines

Authorisation number:

65883/13-07-2021/K-0211501

Date of authorisation status change:

12/07/2021

Reference member state:

France

Procedure number:

FR/V/0415/001

Concerned member states:

Austria Belgium Cyprus Czechia Denmark Estonia Finland Germany Greece
Hungary Ireland Italy Latvia Lithuania Luxembourg Malta Netherlands
Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Source URL: <https://medicines.health.europa.eu/veterinary/600000040445>