

# THERIOS 75 MG CHEWABLE TABLETS FOR CATS

Authorised

- Cefalexin monohydrate

## Product identification

**Medicine name:**

THERIOS 75 MG CHEWABLE TABLETS FOR CATS

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**Active substance:**

Cefalexin monohydrate

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**Target species:**

Cat

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Cefalexin monohydrate

78.90 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Chewable tablet

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01DB01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Austria

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**Package description:**

Cardboard box with 1 blister (Polyvinylchloride / thermo-elast / polyvinylidene chloride - aluminium heat-sealed) of 10 tablets

Cardboard box with 20 blisters (Polyvinylchloride / thermo-elast / polyvinylidene chloride - aluminium heat-sealed) of 10 tablets

Cardboard box with 15 blisters (Polyvinylchloride / thermo-elast / polyvinylidene chloride - aluminium heat-sealed) of 10 tablets

Cardboard box with 10 blisters (Polyvinylchloride / thermo-elast / polyvinylidene chloride - aluminium heat-sealed) of 10 tablets

Cardboard box with 2 blisters (Polyvinylchloride / thermo-elast / polyvinylidene chloride - aluminium heat-sealed) of 10 tablets

Cardboard box with 1 blister (Polyamide / aluminium / polyvinylchloride - aluminium heat-sealed) of 10 tablets

Cardboard box with 2 blisters (Polyamide / aluminium / polyvinylchloride - aluminium heat-sealed) of 10 tablets

Cardboard box with 10 blisters (Polyamide / aluminium / polyvinylchloride - aluminium heat-sealed) of 10 tablets

Cardboard box with 15 blisters (Polyamide / aluminium / polyvinylchloride - aluminium heat-sealed) of 10 tablets

Cardboard box with 20 blisters (Polyamide / aluminium / polyvinylchloride - aluminium heat-sealed) of 10 tablets

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ceva Sante Animale

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**Marketing authorisation date:**

6/08/2010

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**Manufacturing sites for batch release:**

Ceva Sante Animale

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**Responsible authority:**

Austrian Agency For Health And Food Safety

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**Authorisation number:**

8-00892

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**Date of authorisation status change:**

6/08/2010

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**Reference member state:**

France

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**Procedure number:**

FR/V/0213/001

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**Concerned member states:**

Austria Belgium Czechia Denmark Finland Germany Greece Hungary  
Ireland Italy Luxembourg Netherlands Norway Portugal Romania Spain  
Sweden United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Package Leaflet

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

### 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.

### Combined File of all Documents