THERIOS 75 MG CHEWABLE TABLETS FOR CATS

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:
THERIOS 75 MG CHEWABLE TABLETS FOR CATS
Therios 75 mg purutabletti

Active substance:
Cefalexin monohydrate

Target species:
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Cefalexin monohydrate
78.90 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01DB01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Finland

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

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
16/12/2010

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Finnish Medicines Agency

Authorisation number:
27742

Date of authorisation status change:
16/12/2010

Reference member state:
France

Procedure number:
FR/V/0213/001

Concerned member states:
Austria Belgium Czechia Denmark Finland Germany Greece Hungary Ireland Italy Luxembourg Netherlands Norway Portugal Romania 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

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

Package Leaflet

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