THERIOS 75 MG CHEWABLE TABLETS FOR CATS

Not authorised

- Cefalexin monohydrate

Product identification

Medicine name:
THERIOS 75 MG CHEWABLE TABLETS FOR CATS
Therios 75 mg kauwtabletten voor katten

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:
Surrendered

Authorised in:
Netherlands

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 B.V.

Marketing authorisation date:
30/06/2010

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 105591

Date of authorisation status change:
29/10/2025

Reference member state:
France

Procedure number:
FR/V/0213/001

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

Documents

Combined File of all Documents

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