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Therios 750 mg palatable tablets for dogs

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:
Therios 750 mg palatable tablets for dogs

Active substance:
Cefalexin monohydrate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Cefalexin monohydrate
788.86 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:
Greece

Available in:
Greece

Package description:
Polyvinylchloride blister heat sealed with an aluminium cover foil.Pack sizes:Cardboard box with 20 blisters of 10 tablets
Polyvinylchloride blister heat sealed with an aluminium cover foil.Pack sizes:Cardboard box with 3 blisters of 10 tablets
Polyvinylchloride blister heat sealed with an aluminium cover foil.Pack sizes:Cardboard box with 1 blister of 10 tablets

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:
Ceva Hellas LLC

Marketing authorisation date:
30/11/2010

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
National Organization For Medicines

Authorisation number:
16196/21-03-2017/K-0185403

Date of authorisation status change:
20/03/2017

Reference member state:
Ireland

Procedure number:
IE/V/0516/002

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

English (PDF)
Published on: 15/12/2024
Updated on: 14/03/2026
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Combined File of all Documents