File downloaded on 2026-11-16
Source URL: https://medicines.health.europa.eu/veterinary/en/600000032997

Cefatab

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:
Cefatab

Active substance:
Cefalexin monohydrate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Cefalexin monohydrate
1050.00 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:
Netherlands

Package description:
Cardboard box of 9 blisters (Aluminium - PVC /PE /PVDC blister) of 10 tablets
Cardboard box of 8 blisters (Aluminium - PVC /PE /PVDC blister) of 10 tablets
Cardboard box of 7 blisters (Aluminium - PVC /PE /PVDC blister) of 10 tablets
Cardboard box of 6 blisters (Aluminium - PVC /PE /PVDC blister) of 10 tablets
Cardboard box of 5 blisters (Aluminium - PVC /PE /PVDC blister) of 10 tablets
Cardboard box of 4 blisters (Aluminium - PVC /PE /PVDC blister) of 10 tablets
Cardboard box of 3 blisters (Aluminium - PVC/ PE /PVDC blister) of 10 tablets
Cardboard box of 25 blisters (Aluminium - PVC /PE /PVDC blister) of 10 tablets.
Cardboard box of 10 blisters (Aluminium - PVC /PE /PVDC blister) of 10 tablets.
Cardboard box containing 10 separate cardboard boxes, each containing 1 blister of 10 tablets.
Cardboard box of 1 blister (Aluminium - PVC /PE /PVDC blister) of 10 tablets
Cardboard box of 2 blisters (Aluminium - PVC/ PE /PVDC blister) 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:
CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:
4/07/2019

Manufacturing sites for batch release:
CP-Pharma Handelsgesellschaft mbH

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 123299

Date of authorisation status change:
14/02/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0263/004

Concerned member states:
Germany

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

Documents

Combined File of all Documents

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