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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
525.00 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:
Germany

Available in:
Germany

Package description:
Cardboard box of 1 blister (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 2 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 3 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 5 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 9 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 8 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:
23/03/2019

Manufacturing sites for batch release:
CP-Pharma Handelsgesellschaft mbH

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402587.00.00

Date of authorisation status change:
22/01/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0263/003

Concerned member states:
Germany

To consult adverse reactions on veterinary medicinal products please go to 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.