

Cefabactin 500 mg tablets for dogs and cats

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:

Cefabactin 500 mg tablets for dogs and cats

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:

Cyprus

Available in:

Cyprus

Package description:

Cardboard box of 4 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets

Cardboard box of 3 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets

Cardboard box of 6 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets

Cardboard box of 5 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets

Cardboard box of 25 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets

Cardboard box of 2 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets

Cardboard box of 10 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets. (Cardboard box containing 10 separate cardboard boxes, each containing 1 blister of 10 tablets)

Cardboard box of 7 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets

Cardboard box of 9 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets

Cardboard box of 8 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets

Cardboard box of 1 blister (Aluminium - PVC/PE/PVDC) of 10 tablets

Cardboard box of 10 blisters (Aluminium - PVC/PE/PVDC) 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:

Le Vet. Beheer B.V.

Marketing authorisation date:

6/07/2016

Manufacturing sites for batch release:

Lelypharma B.V.

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

CY00559V

Date of authorisation status change:

6/07/2016

Reference member state:

Netherlands

Procedure number:

NL/V/0201/003

Concerned member states:

Austria Belgium Croatia Cyprus Czechia Denmark Estonia Finland France
Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania
Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain
Sweden United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

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