

# Cefabactin 500 mg tablets for dogs and cats

Authorised

- Cefalexin monohydrate

## Product identification

**Medicine name:**

Cefabactin 500 mg tablets for dogs and cats

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**Active substance:**

Cefalexin monohydrate

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Cefalexin monohydrate

525.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01DB01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Denmark

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**Available in:**

Denmark

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Le Vet. Beheer B.V.

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**Marketing authorisation date:**

31/08/2016

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**Manufacturing sites for batch release:**

Lelypharma B.V.

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**Responsible authority:**

Danish Medicines Agency

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**Authorisation number:**

56261

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**Date of authorisation status change:**

31/08/2016

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0201/003

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**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

Summary of Product Characteristics

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