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:
Romania

Available in:
Romania

Package description:
Cardboard box of 1 blister (Aluminium - PVC/PE/PVDC) of 10 tablets
Cardboard box of 8 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets
Cardboard box of 9 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets
Cardboard box of 7 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 2 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets
Cardboard box of 25 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets
Cardboard box of 5 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets
Cardboard box of 6 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets
Cardboard box of 3 blisters (Aluminium - PVC/PE/PVDC) of 10 tablets
Cardboard box of 4 blisters (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:
13/07/2016

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
210130

Date of authorisation status change:
16/03/2026

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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