Cephacare flavour 500 mg tablets for dogs

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:
Cephacare flavour 500 mg tablets for dogs

Active substance:
Cefalexin monohydrate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Cefalexin monohydrate
525.91 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:
Ireland

Available in:
Ireland

Package description:
Cephacare flavour 500 mg tablets are supplied in PVC/aluminium foil blister packs each containing 10 tablets, in cardboardboxes containing 20 tablets.
Cephacare flavour 500 mg tablets are supplied in PVC/aluminium foil blister packs each containing 10 tablets, in cardboardboxes containing 100 tablets.
Cephacare flavour 500 mg tablets are supplied in PVC/aluminium foil blister packs each containing 10 tablets, in cardboardboxes containing 250 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:
Ecuphar

Marketing authorisation date:
19/12/2008

Manufacturing sites for batch release:
Lelypharma B.V.

Produlab Pharma B.V.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10491/005/003

Date of authorisation status change:
19/12/2008

Reference member state:
Ireland

Procedure number:
IE/V/0454/003

Concerned member states:
Austria Belgium France Germany Luxembourg Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents

English (PDF)
Published on: 6/04/2025

Download