

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:

United Kingdom (Northern Ireland)

Available in:

United Kingdom (Northern 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:

The Veterinary Medicines Directorate

Authorisation number:

Vm 32742/4030

Date of authorisation status change:

22/06/2024

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