Cardisure Flavoured 5 mg Tablets for Dogs

Authorised

- Pimobendan

Product identification

Medicine name:
Cardisure Flavoured 5 mg Tablets for Dogs
Cardisure 5 mg Tablet
Cardisure 5 mg Comprimé
Cardisure 5 mg Tablette

Active substance:
Pimobendan

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Pimobendan
5.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC01CE90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
20 blisters per carton, 5 tablets per blister (Aluminium - Aluminium blister)
4 blisters per carton, 5 tablets per blister (Aluminium - Aluminium blister)
25 blisters per carton, 10 tablets per blister (Aluminium - PVC,PE,PVDC blister)
2 blisters per carton, 10 tablets per blister (Aluminium - PVC,PE,PVDC blister)
10 blisters per carton, 5 tablets per blister (Aluminium - Aluminium blister)
10 blisters per carton, 10 tablets per blister (Aluminium - PVC,PE,PVDC blister)
50 blisters per carton, 5 tablets per blister (Aluminium - Aluminium blister)
5 blisters per carton, 10 tablets per blister (Aluminium - PVC,PE,PVDC blister)

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:
Dechra Regulatory B.V.

Marketing authorisation date:
12/07/2011

Manufacturing sites for batch release:
Eurovet Animal Health B.V.
GENERA d.d.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
12/07/2011

Reference member state:
Netherlands

Procedure number:
NL/V/0280/003

Concerned member states:
Austria Belgium Denmark Finland France Germany Greece Ireland Italy Luxembourg Norway Poland Portugal Spain Sweden
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

Labelling

This document does not exist in this language (English). You can find it in another language below.

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