Pimocard 2.5 mg Flavoured Tablets for Dogs

Authorised

- Pimobendan

Product identification

Medicine name:
Pimocard 2.5 mg Flavoured Tablets for Dogs

Active substance:
Pimobendan

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Pimobendan
2.50 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:
Cyprus

Available in:
Cyprus

Package description:
5 Aluminium-aluminium blisters with 10 tablets per blister in a carton box.
2 Aluminium-PVC/PE/PVDC blisters with 10 tablets per blister in a carton box.
5 Aluminium-PVC/PE/PVDC blisters with 10 tablets per blister in a carton box.
25 Aluminium-aluminium blisters with 10 tablets per blister in a carton box.
25 Aluminium-PVC/PE/PVDC blisters with 10 tablets per blister in a carton box.
2 Aluminium-aluminium blisters with 10 tablets per blister in a carton box.
10 Aluminium-aluminium blisters with 10 tablets per blister in a carton box.
10 Aluminium-PVC/PE/PVDC blisters with 10 tablets per blister in a carton box.

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:
24/09/2018

Manufacturing sites for batch release:
Dales Pharmaceuticals Limited
Eurovet Animal Health B.V.
Genera d.d.

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00700V

Date of authorisation status change:
24/09/2018

Reference member state:
Netherlands

Procedure number:
NL/V/0283/002

Concerned member states:
Belgium Bulgaria Croatia Cyprus Czechia Estonia Germany Hungary Ireland Italy Latvia Lithuania Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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