Furosivet 20 mg Tablets for Dogs and Cats

Authorised

- Furosemide

Product identification

Medicine name:
Furosivet 20 mg Tablets for Dogs and Cats

Active substance:
Furosemide

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Furosemide
20.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC03CA01

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

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Package description:
White high-density polyethylene container containing 250 tablets, with a silica gel bag desiccant, and sealed with a white, child-resistant, polypropylene closure.
Cardboard box of 1 Aluminium-PVC foil/PVC-PVDC blister of 10 tablets.
Cardboard box of 2 Aluminium-PVC foil/PVC-PVDC blisters of 10 tablets.
Cardboard box of 3 Aluminium-PVC foil/PVC-PVDC blisters of 10 tablets.
Cardboard box of 4 Aluminium-PVC foil/PVC-PVDC blisters of 10 tablets.
Cardboard box of 5 Aluminium-PVC foil/PVC-PVDC blisters of 10 tablets.
Cardboard box of 6 Aluminium-PVC foil/PVC-PVDC blisters of 10 tablets.
Cardboard box of 7 Aluminium-PVC foil/PVC-PVDC blisters of 10 tablets.
Cardboard box of 8 Aluminium-PVC foil/PVC-PVDC blisters of 10 tablets.
Cardboard box of 9 Aluminium-PVC foil/PVC-PVDC blisters of 10 tablets.
Cardboard box of 10 Aluminium-PVC foil/PVC-PVDC blisters of 10 tablets.
Cardboard box of 12 Aluminium-PVC foil/PVC-PVDC blisters of 10 tablets.
Cardboard box of 15 Aluminium-PVC foil/PVC-PVDC blisters of 10 tablets.
Cardboard box of 25 Aluminium-PVC foil/PVC-PVDC blisters of 10 tablets.
Cardboard box of 50 Aluminium-PVC foil/PVC-PVDC blisters 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:
Millpledge Europe

Marketing authorisation date:
5/03/2019

Manufacturing sites for batch release:
Millpledge Europe

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 61300/3002

Date of authorisation status change:
1/12/2024

Reference member state:
Netherlands

Procedure number:
NL/V/0329/001

Concerned member states:
Belgium France Germany 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