

Furosoral10 mg tablets for dogs and cats

Authorised

- Furosemide

Product identification

Medicine name:

Furosoral10 mg tablets for dogs and cats

Furosoral vet 10 mg tabletes kaķiem un suņiem

Active substance:

Furosemide

Target species:

Dog

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Furosemide

10.00 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:

Latvia

Package description:

Cardboard box 3 Aluminium-PVDC/PE/PVC blisters with 10 tablets each, respectively corresponding to 30 tablets per box.

Cardboard box of 2 Aluminium-PVDC/PE/PVC blisters with 10 tablets each, respectively corresponding to 20 tablets per box.

Cardboard box of 1 Aluminium-PVDC/PE/PVC blister with 10 tablets each, respectively corresponding to 10 tablets per box.

Cardboard box of 100 Aluminium-PVDC/PVC blisters with 10 tablets each. Cardboard box containing 10 separate cardboard boxes, each containing 1 blister of 10 tablets.

Cardboard box of 4 Aluminium-PVDC/PVC blisters with 10 tablets each

Cardboard box of 10 Aluminium-PVDC/PVC blisters with 10 tablets each. Cardboard box containing 10 separate cardboard boxes, each containing 1 blister of 10 tablets.

Cardboard box of 6 Aluminium-PVDC/PVC blisters with 10 tablets each

Cardboard box of 50 Aluminium-PVDC/PVC blisters with 10 tablets each. Cardboard box containing 10 separate cardboard boxes, each containing 1 blister of 10 tablets.

Cardboard box of 5 Aluminium-PVDC/PVC blisters with 10 tablets each

Cardboard box of 9 Aluminium-PVDC/PVC blisters with 10 tablets each

Cardboard box of 8 Aluminium-PVDC/PVC blisters with 10 tablets each

Cardboard box of 7 Aluminium-PVDC/PVC blisters with 10 tablets each

Cardboard box of 25 Aluminium-PVDC/PVC blisters with 10 tablets each. Cardboard box containing 10 separate cardboard boxes, each containing 1 blister of 10 tablets.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Le Vet. Beheer B.V.

Marketing authorisation date:

31/10/2014

Manufacturing sites for batch release:

Artesan Pharma GmbH & Co. KG

Lelypharma B.V.

Genera d.d.

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/MRP/14/0062

Date of authorisation status change:

31/10/2014

Reference member state:

Netherlands

Procedure number:

NL/V/0192/001

Concerned member states:

Austria Belgium Croatia Cyprus Czechia Denmark Estonia Finland France
Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania
Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain

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

Documents

Combined File of all Documents

Labelling

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.

Summary of Product Characteristics

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