

# Furosoral 40 mg tablets for dogs and cats

Authorised

- Furosemide

## Product identification

**Medicine name:**

Furosoral 40 mg tablets for dogs and cats

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**Active substance:**

Furosemide

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**Target species:**

Dog

Cat

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Furosemide

40.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QC03CA01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Sweden

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**Available in:**

Sweden

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**Package description:**

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

Cardboard box of 100 Aluminium-PVDC/PVC blisters with 10 tablets each.

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

Cardboard box of 2 Aluminium-PVDC/PVC blisters with 10 tablets each

Cardboard box of 25 Aluminium-PVDC/PVC blisters with 10 tablets each.

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

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

Cardboard box of 3 Aluminium-PVDC/PVC blisters with 10 tablets each

Cardboard box of 1 Aluminium-PVDC/PVC blister with 10 tablets each

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

Cardboard box of 9 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 1 Aluminium-PVDC/PE/PVC blister with 10 tablets, respectively corresponding to 10 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 3 Aluminium-PVDC/PE/PVC blisters with 10 tablets each, respectively corresponding to 30 tablets per box.

Cardboard box of 4 Aluminium-PVDC/PE/PVC blisters with 10 tablets each, respectively corresponding to 40 tablets per box.

Cardboard box of 5 Aluminium-PVDC/PE/PVC blisters with 10 tablets each, respectively corresponding to 50 tablets per box.

Cardboard box of 6 Aluminium-PVDC/PE/PVC blisters with 10 tablets each, respectively corresponding to 60 tablets per box.

Cardboard box of 7 Aluminium-PVDC/PE/PVC blisters with 10 tablets each, respectively corresponding to 70 tablets per box.

Cardboard box of 8 Aluminium-PVDC/PE/PVC blisters with 10 tablets each, respectively corresponding to 80 tablets per box.

Cardboard box of 9 Aluminium-PVDC/PE/PVC blisters with 10 tablets each, respectively corresponding to 90 tablets per box.

Cardboard box of 10 Aluminium-PVDC/PE/PVC blisters with 10 tablets each, respectively corresponding to 100 tablets per box.

Cardboard box of 25 Aluminium-PVDC/PE/PVC blisters with 10 tablets each, respectively corresponding to 250 tablets per box.

Cardboard box of 50 Aluminium-PVDC/PE/PVC blisters with 10 tablets each, respectively corresponding to 500 tablets per box.

Cardboard box of 100 Aluminium-PVDC/PE/PVC blisters with 10 tablets each, respectively corresponding to 1000 tablets per box.

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## Additional information

### **Entitlement type:**

Marketing Authorisation

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### **Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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### **Marketing authorisation holder:**

Le Vet. Beheer B.V.

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### **Marketing authorisation date:**

13/11/2014

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### **Manufacturing sites for batch release:**

Artesan Pharma GmbH & Co. KG

Lelypharma B.V.

Genera d.d.

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**Responsible authority:**

Swedish Medical Products Agency

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**Authorisation number:**

51335

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**Date of authorisation status change:**

13/11/2014

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0192/002

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**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  
Sweden United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

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.

## Combined File of all Documents

## Labelling

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