Furosoral10 mg tablets for dogs and cats

Authorised

- Furosemide

Product identification

Medicine name:
Furosoral10 mg tablets for dogs and cats
Furosoral 10 mg tabletten voor katten en honden

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:
Netherlands

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:
Ast Farma B.V.

Marketing authorisation date:
14/03/2014

Manufacturing sites for batch release:
Artesan Pharma GmbH & Co. KG
Lelypharma B.V.
Genera d.d.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 113962

Date of authorisation status change:
24/01/2022

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

Sweden United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

English (PDF)
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