Furosoral 40 mg tablets for dogs and cats

Authorised

- Furosemide

Product identification

Medicine name:
Furosoral 40 mg tablets for dogs and cats
Furosoral 40 mg Tabletten für Katzen und Hunde

Active substance:
Furosemide

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

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

Pharmaceutical form:
Tablet

Withdrawal period by route of administration:

Oral use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QC03CA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Germany

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 containing 10 separate cardboard boxes, each containing 1 blister of 10 tablets
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 2 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
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 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 3 Aluminium-PVDC/PVC blisters with 10 tablets each
Cardboard box of 1 Aluminium-PVDC/PVC blister with 10 tablets
Cardboard box of 8 Aluminium-PVDC/PVC blisters with 10 tablets each
Cardboard box of 9 Aluminium-PVDC/PVC blisters with 10 tablets each

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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:
4/12/2019

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

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402694.00.00

Date of authorisation status change:
4/12/2019

Reference member state:
Netherlands

Procedure number:
NL/V/0192/002

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)

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

Documents

Combined File of all Documents

English (PDF)
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