File downloaded on 2026-02-10
Source URL: https://medicines.health.europa.eu/veterinary/en/600000108429

NELIO 2.5 MG TABLET FOR CATS

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:
NELIO 2.5 MG TABLET FOR CATS
Nelio 2,5 mg Tabletten für Katzen

Active substance:
Benazepril hydrochloride

Target species:
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Benazepril hydrochloride
2.50 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QC09AA07

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

Authorisation status:
Valid

Authorised in:
Austria

Package description:
Box with 18 strips (Aluminium/Aluminium) of 10 tablets
Box with 1 strip (Polyamide-Aluminium-Desicant/Aluminium) of 10 tablets
Box with 2 strips (Polyamide-Aluminium-Desicant/Aluminium) of 10 tablets
Box with 5 strips (Polyamide-Aluminium-Desicant/Aluminium) of 10 tablets
Box with 10 strips (Polyamide-Aluminium-Desicant/Aluminium) of 10 tablets
Box with 14 strips (Polyamide-Aluminium-Desicant/Aluminium) of 10 tablets
Box with 18 strips (Polyamide-Aluminium-Desicant/Aluminium) of 10 tablets
Box with 1 strip (Aluminium/Aluminium) of 10 tablets
Box with 14 strips (Aluminium/Aluminium) of 10 tablets
Box with 10 strips (Aluminium/Aluminium) of 10 tablets
Box with 5 strips (Aluminium/Aluminium)of 10 tablets
Box with 2 strips (Aluminium/Aluminium) 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:
Ceva Sante Animale

Marketing authorisation date:
25/02/2010

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-00854

Date of authorisation status change:
25/02/2010

Reference member state:
France

Procedure number:
FR/V/0178/002

Concerned member states:
Austria Belgium Czechia Finland Germany Hungary Ireland Italy Luxembourg Netherlands Poland Portugal Romania Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to 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.

Labelling

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