

# NELIO 5 MG TABLET FOR CATS

Authorised

- Benazepril hydrochloride

## Product identification

**Medicine name:**

NELIO 5 MG TABLET FOR CATS

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**Active substance:**

Benazepril hydrochloride

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**Target species:**

Cat

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Benazepril hydrochloride  
5.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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

QC09AA07

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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:**

France

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**Available in:**

France

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**Package description:**

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Box with 1 strip (Polyamide-Aluminium-Desiccant/Aluminium) of 10 tablets

Box with 2 strips (Polyamide-Aluminium-Desiccant/Aluminium) of 10 tablets

Box with 3 strips (Polyamide-Aluminium-Desiccant/Aluminium) of 10 tablets

Box with 5 strips (Polyamide-Aluminium-Desiccant/Aluminium) of 10 tablets

Box with 10 strips (Polyamide-Aluminium-Desiccant/Aluminium) of 10 tablets

Box with 20 strips (Polyamide-Aluminium-Desiccant/Aluminium) of 10 tablets

Box with 50 strips (Polyamide-Aluminium-Desiccant/Aluminium) of 10 tablets

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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:**

Ceva Sante Animale

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**Marketing authorisation date:**

17/12/2008

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**Manufacturing sites for batch release:**

Ceva Sante Animale

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/5121330 5/2008

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**Date of authorisation status change:**

17/12/2013

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**Reference member state:**

France

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**Procedure number:**

FR/V/0178/001

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**Concerned member states:**

Austria Belgium Czechia Finland Germany Hungary Ireland Italy  
Luxembourg Netherlands Poland Portugal Romania Spain  
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

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

Package Leaflet and Labelling

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