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NELIO 5 MG TABLET FOR DOGS

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:

NELIO 5 MG TABLET FOR DOGS

Active substance:

Benazepril hydrochloride

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Benazepril hydrochloride
5.00 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:

Poland

Available in:

Poland

Package description:

Cardboard box with 1 blister strip (Aluminium/Aluminium) of 10 tablets

Cardboard box with 25 blister strips (Aluminium/Aluminium) of 10 tablets

Cardboard box with 10 blister strips (Aluminium/Aluminium) of 10 tablets

Cardboard box with 5 blister strips (Aluminium/Aluminium) of 10 tablets

Cardboard box with 1 blister strip (Poylamide-Aluminium-Desicant/Aluminium) of 10 tablets

Cardboard box with 5 blisters strip (Poylamide-Aluminium-Desicant/Aluminium) of 10 tablets

Cardboard box with 10 blisters strip (Poylamide-Aluminium-Desicant/Aluminium) of 10 tablets

Cardboard box with 25 blisters strip (Poylamide-Aluminium-Desicant/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 Animal Health Polska Sp. z o.o.

Marketing authorisation date:

15/10/2010

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

2021

Date of authorisation status change:

15/10/2010

Reference member state:

France

Procedure number:

FR/V/0205/001

Concerned member states:

Austria Belgium Denmark Finland Germany Greece 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

Labelling

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.

Package Leaflet

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

Combined File of all Documents