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# Canergy 100 mg Tablets for Dogs

Authorised

- Propentofylline

## Product identification

**Medicine name:**

Canergy 100 mg Tablets for Dogs  
Canergy 100 mg tableta kutya'knak

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**Active substance:**

Propentofylline

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Propentofylline  
100.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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

QC04AD90

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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:**

Hungary

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**Package description:**

Cardboard box with 7 Aluminium - PA\_ALU\_PVC blister of 10 tablets  
Cardboard box with 6 Aluminium - PA\_ALU\_PVC blister of 10 tablets  
Cardboard box with 50 Aluminium - PA\_ALU\_PVC blister of 10 tablets  
Cardboard box with 5 Aluminium - PA\_ALU\_PVC blister of 10 tablets  
Cardboard box with 4 Aluminium - PA\_ALU\_PVC blister of 10 tablets  
Cardboard box with 3 Aluminium - PA\_ALU\_PVC blister of 10 tablets  
Cardboard box with 25 Aluminium - PA\_ALU\_PVC blister of 10 tablets  
Cardboard box with 2 Aluminium - PA\_ALU\_PVC blister of 10 tablets  
Cardboard box with 10 Aluminium - PA\_ALU\_PVC blister of 10 tablets  
Cardboard box with 1 Aluminium - PA\_ALU\_PVC blister of 10 tablets  
Cardboard box with 9 Aluminium - PA\_ALU\_PVC blister of 10 tablets  
Cardboard box with 8 Aluminium - PA\_ALU\_PVC blister 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:**

Le Vet. Beheer B.V.

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**Marketing authorisation date:**

6/05/2015

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**Manufacturing sites for batch release:**

Artesan Pharma GmbH & Co. KG  
Lelypharma B.V.  
Genera d.d.

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**Responsible authority:**

Directorate Of Veterinary Medicinal Products

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**Authorisation number:**

3647/X/15 NÉBIH ÁTI

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**Date of authorisation status change:**

6/05/2015

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0313/001

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**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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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)