

# Apovomin 1 mg/ml solution for injection for dogs

Authorised

- Apomorphine hydrochloride hemihydrate

## Product identification

**Medicine name:**

Apovomin 1 mg/ml solution for injection for dogs

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**Active substance:**

Apomorphine hydrochloride hemihydrate

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**Target species:**

Dog

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Apomorphine hydrochloride hemihydrate  
1.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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

QN04BC07

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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:**

Italy

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**Package description:**

Clear Type I glass vials containing 5 ml, closed with a coated Type I bromobutyl rubber stopper and sealed with an aluminium cap. Each vial is packed into a cardboard box.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Dechra Regulatory B.V.

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**Marketing authorisation date:**

6/05/2021

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

105482

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**Date of authorisation status change:**

6/05/2021

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0343/001

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**Concerned member states:**

Austria Bulgaria Croatia Czechia Denmark Estonia Finland France Greece  
Hungary Iceland Ireland Italy Latvia Lithuania 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)

## Documents

Combined File of all Documents

English (PDF)

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