

# Apovomin 3 mg/ml solution for injection for dogs

Authorised

- Apomorphine hydrochloride hemihydrate

## Product identification

**Medicine name:**

Apovomin 3 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  
3.00 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:**

Spain

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**Available in:**

Spain

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**Package description:**

Clear Type I glass vial containing 10ml, closed with a coated bromobutyl rubber stopper and sealed with analuminium cap. Each vial is packed into a cardboard box. Pack size: Multi-pack with 10 vials of 10 ml

Clear Type I glass vial containing 5ml, closed with a coated bromobutyl rubber stopper and sealed with analuminium cap. Each vial is packed into a cardboard box. Pack size: Multi-pack with 10 vials of 5 ml

Clear Type I glass vial containing 20 ml, closed with a coated bromobutyl rubber stopper and sealed with analuminium cap. Each vial is packed into a cardboard box. Pack size: Box with 1 vial of 20 ml

Clear Type I glass vial containing 10ml, closed with a coated bromobutyl rubber stopper and sealed with analuminium cap. Each vial is packed into a cardboard box. Pack size: Box with 1 vial of 10 ml

Clear Type I glass vial containing 5ml, closed with a coated bromobutyl rubber stopper and sealed with analuminium cap. Each vial is packed into a cardboard box. Pack size: Box with 1 vial of 5 ml

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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:**

Dechra Regulatory B.V.

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**Marketing authorisation date:**

8/02/2019

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

3740 ESP

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**Date of authorisation status change:**

8/02/2019

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0482/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia Finland France  
Germany Greece Hungary Iceland Italy Latvia Lithuania Luxembourg  
Netherlands 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

### Summary of Product Characteristics

English (PDF)

Published on: 16/01/2026

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### Package Leaflet

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.

### Combined File of all Documents