

# PropoFlo Plus, 10mg/ml, Emulsion for Injection

Authorised

- Propofol

## Product identification

**Medicine name:**

PropoFlo Plus, 10mg/ml, Emulsion for Injection

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**Active substance:**

Propofol

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**Target species:**

Dog

Cat

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**Route of administration:**

Intravenous use

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## Product details

**Active substance and strength:**

Propofol

10.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Emulsion for injection

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

QN01AX10

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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:**

Norway

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**Available in:**

Norway

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**Package description:**

box containing 1 vial of 50 ml

box containing 5 vials of 20 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Zoetis Animal Health ApS

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**Marketing authorisation date:**

27/06/2012

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**Manufacturing sites for batch release:**

Fresenius Kabi AB

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**Responsible authority:**

Norwegian Medical Products Agency

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**Authorisation number:**

11-8713

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**Date of authorisation status change:**

27/06/2012

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0324/001

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**Concerned member states:**

Austria Belgium Denmark Finland France Germany Ireland Italy

Luxembourg Netherlands Norway Portugal 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: 6/04/2023

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Package Leaflet

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

Combined File of all Documents

eu-PUAR-propoflo-plus--10mg-ml--emulsion-for-injection-en.pdf