

# Comfortan 10 mg/ml solution for injection for dogs and cats

Authorised

- Methadone hydrochloride

## Product identification

**Medicine name:**

Comfortan 10 mg/ml solution for injection for dogs and cats

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**Active substance:**

Methadone hydrochloride

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**Target species:**

Dog

Cat

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**Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

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## Product details

**Active substance and strength:**

Methadone hydrochloride

10.00 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:**

QN02AC90

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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:**

Denmark

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**Available in:**

Denmark

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**Package description:**

Cardboard box containing 1 vial (uncoloured glass type I, closed teflon-coated chlorobutyl rubber stopper type I, secured with an aluminium cap) filled with 50 ml  
Cardboard box containing 1 vial (uncoloured glass type I, closed teflon-coated chlorobutyl rubber stopper type I, secured with an aluminium cap) filled with 5 ml  
Cardboard box containing 1 vial (uncoloured glass type I, closed teflon-coated chlorobutyl rubber stopper type I, secured with an aluminium cap) filled with 30 ml  
Cardboard box containing 1 vial (uncoloured glass type I, closed teflon-coated chlorobutyl rubber stopper type I, secured with an aluminium cap) filled with 25 ml  
Cardboard box containing 1 vial (uncoloured glass type I, closed teflon-coated chlorobutyl rubber stopper type I, secured with an aluminium cap) filled with 20 ml  
Cardboard box containing 1 vial (uncoloured glass type I, closed teflon-coated chlorobutyl rubber stopper type I, secured with an aluminium cap) filled with 10 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:**

Eurovet Animal Health B.V.

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**Marketing authorisation date:**

15/04/2011

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**Manufacturing sites for batch release:**

Eurovet Animal Health B.V.

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**Responsible authority:**

Danish Medicines Agency

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**Authorisation number:**

46793

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**Date of authorisation status change:**

15/04/2011

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0150/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Denmark Finland France Germany Ireland  
Italy Norway Poland Portugal 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

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

Combined File of all Documents

PuAR Comfortan REG NL 107389 \_updated 20230831.pdf