

# Fentadon 50 microgram/ml, solution for injection for dogs

Authorised

- Fentanyl citrate

## Product identification

**Medicine name:**

Fentadon 50 microgram/ml, solution for injection for dogs

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**Active substance:**

Fentanyl citrate

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**Target species:**

Dog

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**Route of administration:**

Intravenous use

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## Product details

**Active substance and strength:**

Fentanyl citrate

78.50 microgram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection/infusion

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

QN02AB03

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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:**

Sweden

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**Available in:**

Sweden

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**Package description:**

Vial of uncoloured glass type I, filled with 10 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

Vial of uncoloured glass type I, filled with 100 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

Vial of uncoloured glass type I, filled with 50 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

Vial of uncoloured glass type I, filled with 5 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

Vial of uncoloured glass type I, filled with 30 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

Vial of uncoloured glass type I, filled with 25 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

Vial of uncoloured glass type I, filled with 20 ml. Teflon-coated chlorobutyl rubber stoppers type I secured with aluminium caps.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Well-established use application (Article 13a 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:**

21/02/2013

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**Manufacturing sites for batch release:**

Eurovet Animal Health B.V.

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**Responsible authority:**

Swedish Medical Products Agency

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**Authorisation number:**

44985

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**Date of authorisation status change:**

21/02/2013

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0155/001

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**Concerned member states:**

Austria Belgium Denmark France Germany Italy Luxembourg Norway  
Poland Portugal 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

Package Leaflet

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

### Summary of Product Characteristics

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### Combined File of all Documents

English (PDF)

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### Labelling

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