

# Recudon 2.5 mg/ml + 0.125 mg/ml Solution for Injection for Horses and Dogs

Authorised

- Levomethadone hydrochloride
- Fenpipramide hydrochloride

## Product identification

### Medicine name:

Recudon 2.5 mg/ml + 0.125 mg/ml solution for injection for horses and dogs

Recudon 2.5 mg/ml + 0.125 mg/ml Solution for Injection for Horses and Dogs

### Active substance:

Levomethadone hydrochloride

Fenpipramide hydrochloride

### Target species:

Dog

Horse

### Route of administration:

Intravenous use

Intramuscular use

## Product details

### Active substance and strength:

Levomethadone hydrochloride

2.50 milligram(s) / 1.00 millilitre(s)

Fenpipramide hydrochloride

0.13 milligram(s) / 1.00 millilitre(s)

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### Pharmaceutical form:

Solution for injection

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### Withdrawal period by route of administration:

#### Intravenous use:

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#### Horse

- Meat and offal. 3 day

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

QN02AC52

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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:

United Kingdom (Northern Ireland)

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### Package description:

Cardboard box with 1 clear glass (Type I) vial of 50 ml with a coated bromobutyl rubber stopper and aluminium cap.

Cardboard box with 1 clear glass (Type I) vial of 30 ml with a coated bromobutyl rubber stopper and aluminium cap.

Cardboard box with 1 clear glass (Type I) vial of 10 ml with a coated bromobutyl rubber stopper and aluminium cap.

Cardboard box with 1 clear glass (Type I) vial of 10 ml (5 ml in a 10 ml sized vial) with a coated bromobutyl rubber stopper and aluminium cap.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 18 of Regulation (EU) 2019/6)

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**Marketing authorisation holder:**

Alfasan Nederland B.V.

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**Marketing authorisation date:**

21/07/2023

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**Manufacturing sites for batch release:**

Alfasan Nederland B.V.

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**Responsible authority:**

The Veterinary Medicines Directorate

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**Authorisation number:**

Vm 36408/3017

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**Date of authorisation status change:**

21/07/2023

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0384/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania  
Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain

Sweden United Kingdom (Northern Ireland)

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**Generic of:**

600000004401

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)