

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 + 0,125 mg)/ml Roztwór do wstrzykiwań

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)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

-

Horse

- Meat and offal. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02AC52

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

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.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Alfasan Nederland B.V.

Marketing authorisation date:

3/04/2024

Manufacturing sites for batch release:

Alfasan Nederland B.V.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

3307

Date of authorisation status change:

3/04/2024

Reference member state:

Netherlands

Procedure number:

NL/V/0384/001

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)

Generic of:

600000004401

To consult adverse reactions on veterinary medicinal products please go to 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.

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