# Narcostart (Sedastart) 1 mg/ml solution for injection for cats and dogs

Authorised

Medetomidine hydrochloride

# Product identification

# Medicine name:

Narcostart (Sedastart) 1 mg/ml solution for injection for cats and dogs NARCOSTART SOLUTION INJECTABLE POUR CHATS ET CHIENS

## **Active substance:**

Medetomidine hydrochloride

## **Target species:**

Dog Cat

## Route of administration:

Intramuscular use Intravenous use

# **Product details**

## Active substance and strength:

Medetomidine hydrochloride 1.00 milligram(s) / 1.00 millilitre(s)

# **Pharmaceutical form:**

Solution for injection

# Withdrawal period by route of administration:

Intramuscular use:

- . Dog
- . Cat

Intravenous use:

- . Dog
- Cat

**Anatomical therapeutic chemical veterinary (ATCvet) codes:** ON05CM91

QNUJCHJI

# Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

## Authorisation status:

Valid

## Authorised in:

France

## Package description:

 $1 \times 1$  glass vial with 10 ml. Glass (type I) vials closed with bromobutyl rubber closures secured with aluminium crimp caps.

 $5 \times 1$  glass vial with 10 ml. Glass (type I) vials closed with bromobutyl rubber closures secured with aluminium crimp caps.

# Additional information

## **Entitlement type:**

Marketing Authorisation

## Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

## Marketing authorisation holder:

Le Vet. B.V.

#### Marketing authorisation date:

22/03/2010

#### Manufacturing sites for batch release:

Produlab Pharma B.V.

## **Responsible authority:**

National Veterinary Medicines Agency

## Authorisation number:

FR/V/6607082 9/2010

#### Date of authorisation status change:

12/01/2016

## **Reference member state:**

Netherlands

#### **Procedure number:**

NL/V/0138/001

## **Concerned member states:**

Austria Belgium Czechia Denmark Finland France Greece Hungary Iceland Ireland Italy Luxembourg Poland Portugal Slovakia Spain Sweden United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

# Documents

Combined File of all Documents

Summary of Product Characteristics

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

Source URL: https://medicines.health.europa.eu/veterinary/60000037115