

# Narcostop 5 mg/ml, solution for injection for cats and dogs

Authorised

- Atipamezole hydrochloride

## Product identification

**Medicine name:**

Narcostop 5 mg/ml, solution for injection for cats and dogs  
NARCOSTOP SOLUTION INJECTABLE POUR CHATS ET CHIENS

---

**Active substance:**

Atipamezole hydrochloride

---

**Target species:**

Dog  
Cat

---

**Route of administration:**

Intramuscular use

---

## Product details

**Active substance and strength:**

Atipamezole hydrochloride  
5.00 milligram(s) / 1.00 millilitre(s)

---

**Pharmaceutical form:**

Solution for injection

---

**Withdrawal period by route of administration:****Intramuscular use:**

- Dog
  - Cat
- 

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

QV03AB90

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

France

---

**Package description:**

Cardboard box with 5 vials containing 10 ml. Clear glass (type I) vial with bromobutylrubber stopper (type I) containing 10 ml solution for injection.  
Cardboard box with 10 vials containing 10 ml. Clear glass (type I) vial with bromobutylrubber stopper (type I) containing 10 ml solution for injection.  
Cardboard box with 1 vial containing 10 ml. Clear glass (type I) vial with bromobutylrubber stopper (type I) containing 10 ml solution for injection.

---

## 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:**

21/04/2010

---

**Manufacturing sites for batch release:**

Produlab Pharma B.V.

---

**Responsible authority:**

National Veterinary Medicines Agency

---

**Authorisation number:**

FR/V/3467071 6/2010

---

**Date of authorisation status change:**

22/10/2015

---

**Reference member state:**

Netherlands

---

**Procedure number:**

NL/V/0139/001

---

**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](http://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/600000037040>