

Tramadog, 50 mg/ml, solution for injection for dogs

Authorised

- Tramadol hydrochloride

Product identification

Medicine name:

Tramadog, 50 mg/ml, solution for injection for dogs

Active substance:

Tramadol hydrochloride

Target species:

Dog

Route of administration:

Intramuscular use
Intravenous use

Product details

Active substance and strength:

Tramadol hydrochloride
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02AX02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Package description:

Cardboard box of 10 colourless glass ampoules type I of 1 ml.

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:

Domes Pharma

Marketing authorisation date:

9/04/2018

Manufacturing sites for batch release:

Haupt Pharma Livron

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

3642 ESP

Date of authorisation status change:

25/12/2021

Reference member state:

Netherlands

Procedure number:

NL/V/0228/001

Concerned member states:

Austria Belgium France Germany Ireland Italy Luxembourg Portugal Spain
United Kingdom (Northern Ireland)

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.