

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:

Portugal

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:

21/03/2018

Manufacturing sites for batch release:

Haupt Pharma Livron

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

1184/01/18DFVPT

Date of authorisation status change:

26/04/2022

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