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:
Belgium

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:
1/08/2018

Manufacturing sites for batch release:
Haupt Pharma Livron

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V532622

Date of authorisation status change:
1/08/2018

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

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Package Leaflet

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

Labelling

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