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# Tralieve 80 mg chewable tablets for dogs

Authorised

- Tramadol hydrochloride

## Product identification

**Medicine name:**
Tralieve 80 mg chewable tablets for dogs

**Active substance:**
Tramadol hydrochloride

**Target species:**
Dog

**Route of administration:**
Oral use

## Product details

**Active substance and strength:**
Tramadol hydrochloride
80.00 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**
Chewable tablet

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QN02AX02

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Sweden

**Available in:**
Sweden

**Package description:**
Cardboard box of 10 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 1 Aluminium - PVC/PE/PVDC blister of 10 tablets.
Cardboard box containing 10 separate cardboard boxes, each containing 3 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 3 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 2 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 9 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 8 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 7 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 6 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 5 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 4 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 4 Aluminium - PVC/PE/PVDC blisters of 10 tablets.

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Dechra Regulatory B.V.

**Marketing authorisation date:**
29/06/2018

**Manufacturing sites for batch release:**
Lelypharma B.V.
Genera d.d.

**Responsible authority:**
Swedish Medical Products Agency

**Authorisation number:**
56181

**Date of authorisation status change:**
29/06/2018

**Reference member state:**
Netherlands

**Procedure number:**
NL/V/0231/002

**Concerned member states:**
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden 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.