

# Tramadog 50 mg tablet for dogs

Authorised

- Tramadol hydrochloride

## Product identification

**Medicine name:**

Tramadog 50 mg tablet for dogs

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**Active substance:**

Tramadol hydrochloride

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Tramadol hydrochloride  
50.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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

QN02AX02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Netherlands

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**Package description:**

Box with 1 PVC-PVDC / aluminium thermosealed blister containing 10 tablets each.  
Box with 3 PVC-PVDC / aluminium thermosealed blister containing 10 tablets each.  
Box with 6 PVC-PVDC / aluminium thermosealed blister containing 10 tablets each.  
Box with 10 PVC-PVDC / aluminium thermosealed blisters containing 10 tablets each.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Axience

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**Marketing authorisation date:**

18/04/2022

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**Manufacturing sites for batch release:**

Europeenne De Pharmacotechnie Europhartech

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 127462

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**Date of authorisation status change:**

18/04/2022

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0359/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Greece Hungary Ireland Italy Latvia Lithuania  
Luxembourg Malta Norway Poland Portugal Romania Slovakia Slovenia  
Spain Sweden United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
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## Documents

Combined File of all Documents

English (PDF)

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