Tramadog 50 mg tablet for dogs

Authorised

- Tramadol hydrochloride

Product identification

Medicine name:
Tramadog 50 mg tablet for dogs

Active substance:
Tramadol hydrochloride

Target species:
Dog

Route of administration:
Oral use

Product details

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

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02AX02

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

Authorisation status:
Valid

Authorised in:
Norway

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.

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:
Axience

Marketing authorisation date:
20/04/2022

Manufacturing sites for batch release:
Europeenne De Pharmacotechnie Europhartech

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
20-13676

Date of authorisation status change:

20/04/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0359/001

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)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)
Published on: 7/08/2025
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