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Tramvetol 50 mg tablets for dogs

Authorised

- Tramadol hydrochloride

Product identification

Medicine name:
Tramvetol 50 mg tablets 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:
Greece

Available in:
Greece

Package description:
Box of 10 white PVC/PE/PVDC - aluminium blisters of 10 tablets.
Box of 3 white PVC/PE/PVDC - aluminium blisters of 10 tablets.

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:
Virbac

Marketing authorisation date:
30/01/2020

Manufacturing sites for batch release:
Labiana Pharmaceuticals S.L.U

Responsible authority:
National Organization For Medicines

Authorisation number:
124635/21-10-2026/K-0238201

Date of authorisation status change:

20/10/2026

Reference member state:
Netherlands

Procedure number:
NL/V/0267/001

Concerned member states:
Austria Belgium Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Ireland Latvia Lithuania Norway Poland Portugal Romania Slovakia 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