

Cepedol Vet 50 mg chewable tablets for dogs

Authorised

- Tramadol hydrochloride

Product identification

Medicine name:

Cepedol Vet 50 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
50.00 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:

Latvia

Package description:

Cardboard box of 10 tablets. Aluminium-PVC/Aluminium/oPA blisters, containing 10 tablets.

Cardboard box of 30 tablets. Aluminium-PVC/Aluminium/oPA blisters, containing 10 tablets.

Cardboard box of 50 tablets. Aluminium-PVC/Aluminium/oPA blisters, containing 10 tablets.

Cardboard box of 100 tablets. Aluminium-PVC/Aluminium/oPA blisters, containing 10 tablets.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:

CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:

20/02/2024

Manufacturing sites for batch release:

CP-Pharma Handelsgesellschaft mbH

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/DCP/24/0019

Date of authorisation status change:

20/02/2024

Reference member state:

Netherlands

Procedure number:

NL/V/0406/002

Concerned member states:

Austria Belgium Czechia Denmark Estonia Finland France Germany Greece
Hungary Ireland Italy Latvia Lithuania Norway Poland Portugal Slovakia
Spain Sweden United Kingdom (Northern Ireland)

Generic of:

600000004401

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

Documents

Combined File of all Documents

English (PDF)

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