Cepedol Vet 80 mg chewable tablets for dogs

Authorised

- Tramadol hydrochloride

Product identification

Medicine name:
Cepedol Vet 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:
Denmark

Available in:
Denmark

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:
Hybrid application – change in strength (Article 19(1)(a) of Regulation (EU) 2019/6)

Marketing authorisation holder:
CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:
22/03/2024

Manufacturing sites for batch release:
CP-Pharma Handelsgesellschaft mbH

Responsible authority:
Danish Medicines Agency

Authorisation number:
69082

Date of authorisation status change:
22/03/2024

Reference member state:
Netherlands

Procedure number:
NL/V/0406/003

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)

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

Documents

Combined File of all Documents

Final PuAR Tramatab NLV0406001-004DC.pdf