File downloaded on 2026-11-04
Source URL: https://medicines.health.europa.eu/veterinary/en/600000089931

Caniphedrin 50 mg Tabletten für Hunde

Authorised

- Ephedrine hydrochloride

Product identification

Medicine name:
Caniphedrin 50 mg Tabletten für Hunde

Active substance:
Ephedrine hydrochloride

Target species:
Dog

Route of administration:
Oral use

Product details

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

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG04BX90

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

Authorisation status:
Valid

Authorised in:
Denmark

Available in:
Denmark

Package description:
Heat-sealed blister pack, consisting of aluminium foil and a PVC foil with 10 tablets per blister. Package size: Cardboard box containing 10 blisters of 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:
Vetviva Richter GmbH

Marketing authorisation date:
1/07/2022

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Danish Medicines Agency

Authorisation number:

66510

Date of authorisation status change:
1/07/2022

Reference member state:
Austria

Procedure number:
AT/V/0016/002

Concerned member states:
Belgium Bulgaria Croatia Cyprus Czechia Denmark Finland France Greece Hungary Italy Netherlands 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

Summary of Product Characteristics

at-puar-atv0016002-mr-caeniphedrin-en.pdf