Caniphedrin 50 mg Tabletten für Hunde

Authorised

- Ephedrine hydrochloride

Product identification

Medicine name:
Caniphedrin 50 mg Tabletten für Hunde
Caniphedrin 50 mg таблетки за кучета

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

Withdrawal period by route of administration:

Oral use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG04BX90

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Available in:
Bulgaria

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:
21/12/2021

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-3102

Date of authorisation status change:
21/12/2021

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

English (PDF)
Published on: 21/01/2025
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Package Leaflet

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Labelling

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