

Vitofyllin 50 mg film-coated tablets for dogs

Authorised

- Propentofylline

Product identification

Medicine name:

Vitofyllin 50 mg film-coated tablets for dogs
Vitofyllin 50 mg filmom obložena tableta za pse

Active substance:

Propentofylline

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Propentofylline
50.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Film-coated tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QC04AD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Croatia

Package description:

(ID2) 140 Film-coated tablet: unspecified outer container with 10 Blister (polyvinyl chloride; polyvinylidene chloride; aluminium) each with 14 Film-coated tablet

(ID1) 56 Film-coated tablet: unspecified outer container with 4 Blister (polyvinyl chloride; polyvinylidene chloride; aluminium) each with 14 Film-coated tablet

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:

Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Marketing authorisation date:

5/05/2023

Manufacturing sites for batch release:

Artesan Pharma GmbH & Co. KG

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/23-01/307

Date of authorisation status change:

5/05/2023

Reference member state:

Germany

Procedure number:

DE/V/0198/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland
France Greece Hungary Ireland Italy Latvia Lithuania Luxembourg
Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain
Sweden United Kingdom (Northern Ireland)

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www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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Combined File of all Documents