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 apvalkotās tabletes suņiem

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:
Latvia

Package description:
Available only in Latvian
Available only in Latvian

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:
28/03/2023

Manufacturing sites for batch release:
Artesan Pharma GmbH & Co. KG

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/SRP/23/0014

Date of authorisation status change:

28/03/2023

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)

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

Documents

Combined File of all Documents

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