Vitofyllin 50 mg film-coated tablets for dogs

Authorised

- Propentofylline

Product identification

Medicine name:
Vitofyllin 50 mg film-coated tablets for dogs

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:
Hungary

Available in:
Hungary

Package description:
(ID1) 56 Film-coated tablet: unspecified outer container with 4 Blister (polyvinyl chloride; polyvinylidene chloride; aluminium) each with 14 Film-coated tablet
(ID2) 140 Film-coated tablet: unspecified outer container with 10 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:
2/05/2012

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

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
2/05/2012

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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