Vivitonin 50 mg film-coated tablets

Authorised

- Propentofylline

Product identification

Medicine name:
Vivitonin 50 mg film-coated tablets

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

Withdrawal period by route of administration:
Oral use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC04AD90

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

Authorisation status:
Valid

Authorised in:
Ireland

Package description:
Blister package: PVC 250µm/Aluminium foil 20µm.One pack contains 2 blister strips of 30 tablets

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet (Ireland) Limited

Marketing authorisation date:
29/10/1999

Manufacturing sites for batch release:
Intervet Ges.m.b.H.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10996/127/001

Date of authorisation status change:
29/10/1999

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

Documents

Summary of Product Characteristics