KARSIVAN 50 mg, compresse per cani

Authorised

- Propentofylline

Product identification

Medicine name:
KARSIVAN 50 mg, compresse per cani

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:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC04AD90

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

Authorisation status:
Valid

Authorised in:
Italy

Available in:
Italy

Package description:
Available only in Italian

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 International B.V.

Marketing authorisation date:
30/04/1992

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

Responsible authority:
Ministry Of Health

Authorisation number:
101397

Date of authorisation status change:
28/05/2007

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

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.