

Vidalta 10 mg prolonged-release tablets for cats

Authorised

- Carbimazole

Product identification

Medicine name:

Vidalta 10 mg prolonged-release tablets for cats

VIDALTA 10 MG COMPRIMES A LIBERATION PROLONGEE POUR CHATS

Active substance:

Carbimazole

Target species:

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Carbimazole

10.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH03BB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

High density polyethylene container of 100 tablets closed with polypropylene tamper-evident, child-resistant, screw cap bearing a desiccant.

High density polyethylene container of 30 tablets closed with polypropylene tamper-evident, child-resistant, screw cap bearing a desiccant.

6 high density polyethylene containers of 30 tablets closed with polypropylene tamper-evident, child-resistant, screw cap bearing a desiccant.

6 high density polyethylene container of 100 tablets closed with polypropylene tamper-evident, child-resistant, screw cap bearing a desiccant.

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:

13/12/2011

Manufacturing sites for batch release:

Intervet Ges.m.b.H.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/1408206 5/2011

Date of authorisation status change:

6/01/2017

Reference member state:

Ireland

Procedure number:

IE/V/0442/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Denmark France Germany Greece
Hungary Italy Luxembourg Netherlands Norway Poland Portugal Romania
Spain 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

Package Leaflet and Labelling

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

Summary of Product Characteristics

English (PDF)

Published on: 16/01/2026

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