

# 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

**Withdrawal period by route of administration:**

**Oral use:**

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**Cat**

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QH03BB01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

France

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

13/12/2011

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**Manufacturing sites for batch release:**

Intervet Ges.m.b.H.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/1408206 5/2011

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**Date of authorisation status change:**

6/01/2017

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0442/001

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**Concerned member states:**

Austria Belgium Bulgaria Cyprus Denmark France Germany Greece  
Hungary Italy Luxembourg Netherlands Norway Poland Portugal Romania  
Spain United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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

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

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