# Vidalta 15 mg prolonged-release tablets for cats

Authorised

Carbimazole

## Product identification

#### **Medicine name:**

Vidalta 15 mg prolonged-release tablets for cats Vidalta 15 mg prolonged-release tablets for cats

#### **Active substance:**

Carbimazole

#### **Target species:**

Cat

#### **Route of administration:**

Oral use

## **Product details**

## **Active substance and strength:**

Carbimazole 15.00 milligram(s) / 1.00 Tablet

#### **Pharmaceutical form:**

**Tablet** 

## Withdrawal period by route of administration:

#### Oral use:

. Cat

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

QH03BB01

#### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

Ireland

#### Package description:

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

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

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

6 High density polyethylene containers 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 (Ireland) Limited

## Marketing authorisation date:

9/12/2011

#### Manufacturing sites for batch release:

Intervet Ges.m.b.H.

## **Responsible authority:**

Health Products Regulatory Authority

#### **Authorisation number:**

VPA10996/263/002

### Date of authorisation status change:

9/12/2011

#### **Reference member state:**

Ireland

#### **Procedure number:**

IE/V/0442/002

#### **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

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