

# Veloxa Chewable Tablets for Dogs

Authorised

- Febantel
- Pyrantel
- Praziquantel

## Product identification

**Medicine name:**

Veloxa Chewable Tablets for Dogs

VELOXA COMPRIMES A CROQUER POUR CHIENS

**Active substance:**

Febantel

Pyrantel

Praziquantel

**Target species:**

Dog

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Febantel

150.00 milligram(s) / 1.00 Tablet

Pyrantel  
50.00 milligram(s) / 1.00 Tablet

Praziquantel  
50.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Chewable tablet

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

QP52AA51

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription except for some pack sizes

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**Authorisation status:**

Valid

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**Authorised in:**

France

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**Package description:**

PVC/Aluminium/Polyamide blister-forming laminate with aluminium lidding foil containing 2 or 8 chewable tablets. Box containing 52 blister strips of 2 chewable tablets (104 chewable tablets)

PVC/Aluminium/Polyamide blister-forming laminate with aluminium lidding foil containing 2 or 8 chewable tablets. Box containing 13 blister strips of 8 chewable tablets (104 chewable tablets)

PVC/Aluminium/Polyamide blister-forming laminate with aluminium lidding foil containing 2 or 8 chewable tablets. Box containing 1 blister strip of 2 chewable tablets (2 chewable tablets)

PVC/Aluminium/Polyamide blister-forming laminate with aluminium lidding foil containing 2 or 8 chewable tablets. Box containing 2 blister strips of 2 chewable tablets (4 chewable tablets)

PVC/Aluminium/Polyamide blister-forming laminate with aluminium lidding foil containing 2 or 8 chewable tablets. Box containing 1 blister strips of 8 chewable tablets (8 chewable tablets)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Lavet Kft.

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**Marketing authorisation date:**

28/09/2012

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**Manufacturing sites for batch release:**

Lavet Kft.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/2881077 5/2012

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**Date of authorisation status change:**

16/06/2017

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**Reference member state:**

Hungary

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**Procedure number:**

HU/V/0116/001

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**Concerned member states:**

Austria Belgium Finland France Germany Greece Ireland Italy Luxembourg  
Netherlands Norway Portugal Spain Sweden  
United Kingdom (Northern Ireland)

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[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Summary of Product Characteristics

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

### Package Leaflet and Labelling

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