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# Endogard Plus Flavour Tablets for dogs

Authorised

- Febantel
- Praziquantel
- Pyrantel embonate

## Product identification

**Medicine name:**

Endogard Plus Flavour Tablets for dogs

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**Active substance:**

Febantel

Praziquantel

Pyrantel embonate

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Febantel

150.00 milligram(s) / 1.00 Tablet

Praziquantel

50.00 milligram(s) / 1.00 Tablet

Pyrantel embonate

144.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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

QP52AC55

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**Legal status of supply:**

Veterinary medicinal product not subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Netherlands

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**Available in:**

Netherlands

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**Package description:**

Nature of container: Print and perforated Alu-Alu blister: 300 tablets (30 blisters with 10 tablets), in a box.

Nature of container: Print and perforated Alu-Alu blister: 100 tablets (10 blisters with 10 tablets), in a box.

Nature of container: Print and perforated Alu-Alu blister: 50 tablets (5 blisters with 10 tablets), in a box.

Nature of container: Print and perforated Alu-Alu blister: 30 tablets (3 blisters with 10 tablets), in a box.

Nature of container: Print and perforated Alu-Alu blister: 10 tablets (1 blister with 10 tablets), in a box.

Nature of container: Print and perforated Alu-Alu blister: 4 tablets (2 blisters with 2 tablets), in a box.

Nature of container: Print and perforated Alu-Alu blister: 2 tablets (1 blister with 2 tablets), in a box.

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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:**

KRKA tovarna zdravil d.d. Novo mesto

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**Marketing authorisation date:**

21/01/2011

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**Manufacturing sites for batch release:**

KRKA tovarna zdravil d.d. Novo mesto

Krka-Farma d.o.o.

Virbac

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 104801

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**Date of authorisation status change:**

27/01/2022

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0539/001

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**Concerned member states:**

Austria Belgium Czechia Denmark Estonia Germany Greece Hungary Italy  
Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia  
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

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

Summary of Product Characteristics