

# Drontal Dog Tasty Bone

## 150/144/50 mg tablets

Authorised

- Praziquantel
- Pyrantel embonate
- Febantel

## Product identification

**Medicine name:**

Drontal Dog Tasty Bone 150/144/50 mg tablets

---

**Active substance:**

Praziquantel  
Pyrantel embonate  
Febantel

---

**Target species:**

Dog

---

**Route of administration:**

Oral use

---

## Product details

**Active substance and strength:**

Praziquantel

50.00 milligram(s) / 1.00 Tablet

Pyrantel embonate

144.00 milligram(s) / 1.00 Tablet

Febantel

150.00 milligram(s) / 1.00 Tablet

---

**Pharmaceutical form:**

Tablet

---

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

QP52AC55

---

**Legal status of supply:**

Veterinary medicinal product not subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Netherlands

---

**Available in:**

Netherlands

---

**Package description:**

Container material: Blisters formed from PA/Alu/PE foil and sealed with Alu/PE foil. Container size : Cartons containing 2 tablets

Container material: Blisters formed from PA/Alu/PE foil and sealed with Alu/PE foil. Container size : Cartons containing 4 tablets

Container material: Blisters formed from PA/Alu/PE foil and sealed with Alu/PE foil. Container size : Cartons containing 6 tablets

Container material: Blisters formed from PA/Alu/PE foil and sealed with Alu/PE foil. Container size : Cartons containing 24 tablets

Container material: Blisters formed from PA/Alu/PE foil and sealed with Alu/PE foil. Container size : Cartons containing 102 tablets

Container material: Blisters formed from PA/Alu/PE foil and sealed with Alu/PE foil. Container size : Cartons containing 312 tablets

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

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

---

**Marketing authorisation holder:**

Vetoquinol S.A.

---

**Marketing authorisation date:**

11/09/2014

---

**Manufacturing sites for batch release:**

Europeenne De Pharmacotechnie Europhartech  
KVP Pharma+Veterinaer Produkte GmbH

---

**Responsible authority:**

Medicines Evaluation Board

---

**Authorisation number:**

REG NL 114213

---

**Date of authorisation status change:**

26/01/2022

---

**Reference member state:**

Ireland

---

**Procedure number:**

IE/V/0335/001

---

**Concerned member states:**

Austria Denmark Finland France Germany Iceland Italy Netherlands  
Norway Sweden United Kingdom (Northern Ireland)

---

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