

Drontal Tasty Bone Multi-worm XL

525/504/175 mg tablets

Authorised

- Praziquantel
- Pyrantel embonate
- Febantel

Product identification

Medicine name:

DRONTAL DOG FLAVOUR XL (525+504+175)MG/TAB ΔΙΣΚΙΟ

Drontal Tasty Bone Multi-worm XL 525/504/175 mg tablets

Active substance:

Praziquantel

Pyrantel embonate

Febantel

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Praziquantel

175.00 milligram(s) / 1.00 Tablet

Pyrantel embonate
504.00 milligram(s) / 1.00 Tablet
Febantel
525.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Withdrawal period by route of administration:

Oral use:

-

Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AA51

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

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 8 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 48 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:

29/11/2017

Manufacturing sites for batch release:

Europeenne De Pharmacotechnie Europhartech
KVP Pharma + Veterinär Produkte GmbH

Responsible authority:

National Organization For Medicines

Authorisation number:

103783/04-10-2022/K-0205402

Date of authorisation status change:

3/10/2022

Reference member state:

Ireland

Procedure number:

IE/V/0337/002

Concerned member states:

Bulgaria Croatia Cyprus Estonia Greece Hungary Latvia Lithuania Poland
Portugal Romania Slovenia Spain

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

Source URL: <https://medicines.health.europa.eu/veterinary/600000050536>