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

Authorised

- Febantel
- Pyrantel embonate
- Praziquantel

Product identification

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

Active substance:
Febantel

Pyrantel embonate

Praziquantel

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Febantel

525.00 milligram(s) / 1.00 Tablet

Pyrantel embonate
504.00 milligram(s) / 1.00 Tablet

Praziquantel
175.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AA51

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Romania

Available in:
Romania

Package description:
Container material: Blisters formed from PA/Alu/PE foil and sealed with Alu/PE foil.
Container size: Cartons containing 48 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 8 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 2 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:
21/08/2017

Manufacturing sites for batch release:
KVP Pharma+Veterinaer Produkte GmbH
Europeenne De Pharmacotechnie Europhartech

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
220151

Date of authorisation status change:
14/07/2025

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

Combined File of all Documents

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