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

Authorised

- Febantel
- Praziquantel
- Pyrantel embonate

Product identification

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

Active substance:
Febantel

Praziquantel

Pyrantel embonate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:

Febantel
525.00 milligram(s) / 1.00 Tablet

Praziquantel
175.00 milligram(s) / 1.00 Tablet

Pyrantel embonate
504.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AA51

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Cyprus

Available in:
Cyprus

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:
16/05/2018

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

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00687V

Date of authorisation status change:
16/05/2018

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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