

Drontal Tasty Bone Multi-worm 150/144/50 mg tablets

Authorised

- Febantel
- Praziquantel
- Pyrantel embonate

Product identification

Medicine name:

Drontal Tasty Bone Multi-worm 150/144/50 mg tablets

Active substance:

Febantel

Praziquantel

Pyrantel embonate

Target species:

Dog

Route of administration:

Oral use

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

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:

Estonia

Available in:

Estonia

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:

18/09/2014

Manufacturing sites for batch release:

Europeenne De Pharmacotechnie Europhartech
KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:

State Agency Of Medicines

Authorisation number:

1865

Date of authorisation status change:

18/09/2014

Reference member state:

Ireland

Procedure number:

IE/V/0337/001

Concerned member states:

Bulgaria Croatia Cyprus Denmark Estonia Greece Hungary Latvia Lithuania
Netherlands 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

English (PDF)

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