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
Drontal Dog Flavour 150/144/50 mg, tabletės

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

Withdrawal period by route of administration:
Oral use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AC55

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

Authorisation status:
Valid

Authorised in:
Lithuania

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:
2/11/2014

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

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/14/2248/001-006

Date of authorisation status change:
27/10/2019

Reference member state:
Ireland

Procedure number:
IE/V/0337/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia France Greece Hungary Iceland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden

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

Documents

Summary of Product Characteristics

RV2248.pdf
