Drontal Dog Tasty Bone XL 525/504/175 mg tablets

Authorised

- Febantel
- Praziquantel
- Pyrantel embonate

Product identification

Medicine name:
Drontal Dog Tasty Bone XL 525/504/175 mg tablets
Drontal Large Dog Tasty 525/504/175 mg tabletten

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 not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Netherlands

Available in:
Netherlands

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:
2/10/2017

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

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 119735

Date of authorisation status change:
26/01/2022

Reference member state:
Ireland

Procedure number:
IE/V/0335/002

Concerned member states:
Austria Denmark Finland France Germany Iceland Italy Netherlands Norway Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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Summary of Product Characteristics