

Drontal Dog Tasty Bone 150/144/50 mg tablets

Authorised

- Praziquantel
- Pyrantel embonate
- Febantel

Product identification

Medicine name:

Drontal Dog Tasty Bone 150/144/50 mg tablets

Active substance:

Praziquantel
Pyrantel embonate
Febantel

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Praziquantel

50.00 milligram(s) / 1.00 Tablet

Pyrantel embonate

144.00 milligram(s) / 1.00 Tablet

Febantel

150.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AC55

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Iceland

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:

30/08/2014

Manufacturing sites for batch release:

Europeenne De Pharmacotechnie Europhartech
KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:

Icelandic Medicines Agency

Authorisation number:

IS/2/14/008/01

Date of authorisation status change:

9/09/2019

Reference member state:

Ireland

Procedure number:

IE/V/0335/001

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

Summary of Product Characteristics

English (PDF)

Published on: 25/09/2024

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Labelling

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Package Leaflet

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