

Drontal Tasty Bone Wormer XL

525/504/175 mg tablets

Authorised

- Febantel
- Praziquantel
- Pyrantel embonate

Product identification

Medicine name:

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

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 except for some pack sizes

Authorisation status:

Valid

Authorised in:

Czechia

Available in:

Czechia

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:

14/12/2017

Manufacturing sites for batch release:

Europeenne De Pharmacotechnie Europhartech
KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/053/17-C

Date of authorisation status change:

14/12/2017

Reference member state:

Ireland

Procedure number:

IE/V/0338/002

Concerned member states:

Belgium Czechia Luxembourg Slovakia

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 26/08/2024

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Package Leaflet

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Labelling

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