

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

Drontal Plus Tasty 150/144/50 mg Tabletten für Hunde

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

Withdrawal period by route of administration:

Oral use:

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Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AC55

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

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:

14/08/2014

Manufacturing sites for batch release:

KVP Pharma+Veterinaer Produkte GmbH
Europeenne De Pharmacotechnie Europhartech

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

402045.00.00

Date of authorisation status change:

15/05/2020

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

Combined File of all Documents

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