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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:
Germany

Available in:
Germany

Package description:
Container material: Blisters formed from PA/Alu/PE foil and sealed with Alu/PE foil.Container size : Cartons containing 312 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 24 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 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:
14/08/2014

Manufacturing sites for batch release:
KVP Pharma+Veterinar 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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