Drontal Oral Suspension for Puppies 15/5 mg/ml

Authorised

- Febantel
- Pyrantel embonate

Product identification

Medicine name:
Drontal Oral Suspension for Puppies 15/5 mg/ml
Welpan vet. 15 mg/ml + 5 mg/ml mikstur, suspensjon

Active substance:
Febantel
Pyrantel embonate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Febantel
15.00 milligram(s) / 1.00 millilitre(s)

Pyrantel embonate
14.40 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:
Oral use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AF02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Norway

Package description:
Material of the primary container:White high density polyethylene bottle.White polypropylene screw closure.Colourless low density polyethylene adapter insert.Container volume:100ml.Devices supplied (if relevant):5ml transparent polypropylene syringe with rubber plunger.
Material of the primary container:White high density polyethylene bottle.White polypropylene screw closure.Colourless low density polyethylene adapter insert.Container volume:50ml.Devices supplied (if relevant):5ml transparent polypropylene syringe with rubber plunger.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetoquinol S.A.

Marketing authorisation date:
25/03/2009

Manufacturing sites for batch release:
KVP Pharma+Veterinaer Produkte GmbH
Vetoquinol Biowet Sp. z o.o.

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
07-5484

Date of authorisation status change:
22/10/2020

Reference member state:
Ireland

Procedure number:
IE/V/0473/001

Concerned member states:
Austria Estonia Finland France Germany Iceland Latvia Lithuania Norway Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents