

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

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AF02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

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:

6/05/2008

Manufacturing sites for batch release:

KVP Pharma+Veterinaer Produkte GmbH

Vetoquinol Biowet Sp. z o.o.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/1134599 0/2008

Date of authorisation status change:

6/05/2013

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

English (PDF)

Published on: 16/01/2026

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Combined File of all Documents

Package Leaflet and Labelling

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