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# REFORDOG 400 mg/2000 mg spot-on solution for dogs over 25 kg up to 40 kg



- Imidacloprid
- Permethrin (40:60)

# Product identification

#### **Medicine name:**

REFORDOG 400 mg/2000 mg spot-on solution for dogs over 25 kg up to 40 kg REFORDOG 400 MG/2000 MG SOLUTION POUR SPOT-ON POUR CHIENS DE PLUS DE 25 KG A 40 KG

#### **Active substance:**

**Imidacloprid** 

Permethrin (40:60)

### **Target species:**

Dog

#### Route of administration:

Spot-on use

# **Product details**

# **Active substance and strength:**

Imidacloprid 400.00 milligram(s) / 1.00 Pipette Permethrin (40:60) 2000.00 milligram(s) / 1.00 Pipette

#### **Pharmaceutical form:**

Spot-on solution

### **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QP53AC54

### Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

### **Authorisation status:**

Valid

#### Authorised in:

France

### Package description:

Cardboard box containing 1 pipette of 4.0 ml with pouch

Cardboard box containing 2 pipettes of 4.0 ml with pouch

Cardboard box containing 3 pipettes of 4.0 ml with pouch

Cardboard box containing 4 pipettes of 4.0 ml with pouch

Cardboard box containing 6 pipettes of 4.0 ml with pouch

Cardboard box containing 24 pipettes of 4.0 ml with pouch

# Additional information

## **Entitlement type:**

Marketing Authorisation

## Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

# Marketing authorisation holder:

Vetpharma Animal Health S.L.

### Marketing authorisation date:

29/10/2024

# Manufacturing sites for batch release:

Ab7 Sante

### Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

#### **Authorisation number:**

FR/V/9817875 1/2024

### Date of authorisation status change:

29/10/2024

#### Reference member state:

Ireland

#### **Procedure number:**

IE/V/0666/004

#### Concerned member states:

Austria Belgium France Germany Greece Hungary Italy Netherlands Poland Portugal Spain United Kingdom (Northern Ireland)

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

### **Documents**

Package Leaflet and Labelling

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

Summary of Product Characteristics
English (PDF) Published on: 30/11/2025
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Combined File of all Documents