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# REFORDOG 600 mg/3000 mg spot-on solution for dogs over 40 kg up to 60 kg



- Imidacloprid
- Permethrin (40:60)

# Product identification

#### **Medicine name:**

REFORDOG 600 mg/3000 mg spot-on solution for dogs over 40 kg up to 60 kg REFORDOG 600 MG/3000 MG SOLUTION POUR SPOT-ON POUR CHIENS DE PLUS DE 40 KG A 60 KG

#### **Active substance:**

**Imidacloprid** 

Permethrin (40:60)

### **Target species:**

Dog

#### Route of administration:

Spot-on use

# **Product details**

# **Active substance and strength:**

Imidacloprid 600.00 milligram(s) / 1.00 Pipette Permethrin (40:60) 3000.00 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 6.0 ml with pouch

Cardboard box containing 2 pipettes of 6.0 ml with pouch

Cardboard box containing 3 pipettes of 6.0 ml with pouch

Cardboard box containing 4 pipettes of 6.0 ml with pouch

Cardboard box containing 6 pipettes of 6.0 ml with pouch

Cardboard box containing 24 pipettes of 6.0 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/7711657 0/2024

### Date of authorisation status change:

29/10/2024

#### **Reference member state:**

Ireland

#### **Procedure number:**

IE/V/0666/005

#### **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**

Summary of Product Characteristics

English (PDF)

Published on: 30/11/2025

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