

REFORDOG 250 mg/1250 mg spot-on solution for dogs over 10 kg up to 25 kg

Authorised

- Imidacloprid
- Permethrin (40:60)

Product identification

Medicine name:

REFORDOG 250 mg/1250 mg spot-on solution for dogs over 10 kg up to 25 kg

Refordog 250 mg/1250 mg solution pour spot-on pour chiens de plus de 10 kg à 25 kg

Refordog 250 mg/1250 mg spot-on oplossing voor honden vanaf 10 kg tot 25 kg

Refordog 250 mg/1250 mg Lösung zum Auftropfen für Hunde über 10 kg bis 25 kg

Active substance:

Imidacloprid

Permethrin (40:60)

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Imidacloprid

250.00 milligram(s) / 1.00 Pipette

Permethrin (40:60)

1250.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AC54

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Package description:

Cardboard box containing 1 pipette of 2.5 ml with pouch

Cardboard box containing 2 pipettes of 2.5 ml with pouch

Cardboard box containing 3 pipettes of 2.5 ml with pouch

Cardboard box containing 6 pipettes of 2.5 ml with pouch

Cardboard box containing 24 pipettes of 2.5 ml with pouch

Cardboard box containing 4 pipettes of 2.5 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:

10/03/2025

Manufacturing sites for batch release:

Ab7 Sante

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V663998

Date of authorisation status change:

10/03/2025

Reference member state:

Ireland

Procedure number:

IE/V/0666/003

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

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

Labelling

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Summary of Product Characteristics

English (PDF)

Published on: 30/11/2025

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