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 soluzione spot-on per cani di peso superiore a 10 kg e fino a 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

Withdrawal period by route of administration:
Spot-on use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AC54

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Cardboard box containing 1 pipette of 2.5 ml with pouch
Cardboard box containing 1 pipette of 2.5 ml without pouch
Cardboard box containing 2 pipettes of 2.5 ml with pouch
Cardboard box containing 2 pipettes of 2.5 ml without pouch
Cardboard box containing 3 pipettes of 2.5 ml with pouch
Cardboard box containing 3 pipettes of 2.5 ml without pouch
Cardboard box containing 4 pipettes of 2.5 ml without 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 24 pipettes of 2.5 ml without pouch
Cardboard box containing 4 pipettes of 2.5 ml with pouch
Cardboard box containing 6 pipettes of 2.5 ml without 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:
10/01/2025

Manufacturing sites for batch release:
Ab7 Sante

Responsible authority:
European Medicines Agency

Authorisation number:
105649

Date of authorisation status change:
10/01/2025

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