

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

Authorised

- 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 soluzione spot-on per cani di peso superiore a 25 kg e
fino 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

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 4.0 ml without pouch

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 2 pipettes of 4.0 ml without pouch

Cardboard box containing 3 pipettes of 4.0 ml without 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 4 pipettes of 4.0 ml without pouch

Cardboard box containing 6 pipettes of 4.0 ml without pouch

Cardboard box containing 6 pipettes of 4.0 ml with pouch

Cardboard box containing 24 pipettes of 4.0 ml with pouch

Cardboard box containing 24 pipettes of 4.0 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/0666/004

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

Source URL: <https://medicines.health.europa.eu/veterinary/700000131554>