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# PROTECTIX 400 MG/2000 MG SPOT-ON SOLUTION FOR DOGS OVER 25 KG UP TO 40 KG

Authorised

- Imidacloprid
- Permethrin

## Product identification

**Medicine name:**

PROTECTIX 400 MG/2000 MG SPOT-ON SOLUTION FOR DOGS OVER 25 KG UP TO 40 KG

**Active substance:**

Imidacloprid  
Permethrin

**Target species:**

Dog

**Route of administration:**

Cutaneous use

## Product details

**Active substance and strength:**

Imidacloprid

400.00 milligram(s) / 1.00 Pipette

Permethrin

2000.00 milligram(s) / 1.00 Pipette

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**Pharmaceutical form:**

Spot-on solution

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

QP53AC54

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**Legal status of supply:**

Veterinary medicinal product not subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Hungary

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**Package description:**

Pack containing 1 PET/PE/aluminium/surlyn sachet containing one white polypropylene unit dose pipette

Pack containing 2 PET/PE/aluminium/surlyn sachets each containing one white polypropylene unit dose pipette

Pack containing 3 PET/PE/aluminium/surlyn sachets each containing one white polypropylene unit dose pipette

Pack containing 4 PET/PE/aluminium/surlyn sachets each containing one white polypropylene unit dose pipette

Pack containing 6 PET/PE/aluminium/surlyn sachets each containing one white polypropylene unit dose pipette

Pack containing 12 PET/PE/aluminium/surlyn sachets each containing one white polypropylene unit dose pipette

Pack containing 24 PET/PE/aluminium/surlyn sachets each containing one white polypropylene unit dose pipette

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Beaphar B.V.

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**Marketing authorisation date:**

18/12/2023

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**Manufacturing sites for batch release:**

Beaphar B.V.

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**Responsible authority:**

Directorate Of Veterinary Medicinal Products

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**Authorisation number:**

4418/x/23 NÉBIH ÁTI

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**Date of authorisation status change:**

18/12/2023

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**Reference member state:**

France

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**Procedure number:**

FR/V/0433/004

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**Concerned member states:**

Croatia Cyprus Czechia Germany Greece Hungary Italy Latvia Lithuania  
Malta Netherlands Poland Portugal Romania Slovakia Slovenia Spain

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Labelling

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

### Package Leaflet

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

### Summary of Product Characteristics

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

### Combined File of all Documents

eu-puar-frv0433004-mr-rpe805-en.pdf