

# PULSIX 100 mg/500 mg spot-on solution for dogs over 4 kg up to 10 kg

Authorised

- Imidacloprid
- Permethrin (40:60)

## Product identification

### Medicine name:

PULSIX 100 mg/500 mg spot-on solution for dogs over 4 kg up to 10 kg

PULSIX 100 mg + 500 mg Lösung zum Auftropfen für Hunde über 4 kg bis 10 kg

### Active substance:

Imidacloprid

Permethrin (40:60)

### Target species:

Dog

### Route of administration:

Spot-on use

## Product details

### Active substance and strength:

Imidacloprid

100.00 milligram(s) / 1.00 Pipette

Permethrin (40:60)

500.00 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:**

Germany

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**Package description:**

Cardboard box containing 1 pipette of 1.0 ml with pouch

HISTORICAL Cardboard box containing 1 pipette of 1.0 ml without pouch

HISTORICAL Cardboard box containing 2 pipettes of 1.0 ml without pouch

Cardboard box containing 2 pipettes of 1.0 ml with pouch

Cardboard box containing 3 pipettes of 1.0 ml with pouch

HISTORICAL Cardboard box containing 3 pipettes of 1.0 ml without pouch

Cardboard box containing 4 pipettes of 1.0 ml with pouch

HISTORICAL Cardboard box containing 4 pipettes of 1.0 ml without pouch

Cardboard box containing 6 pipettes of 1.0 ml with pouch

HISTORICAL Cardboard box containing 6 pipettes of 1.0 ml without pouch

Cardboard box containing 24 pipettes of 1.0 ml with pouch

HISTORICAL Cardboard box containing 24 pipettes of 1.0 ml without pouch

HISTORICAL Cardboard box containing 12 pipettes of 1.0 ml without pouch

Cardboard box containing 12 pipettes of 1.0 ml with pouch

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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:**

Ab7 Sante

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### **Marketing authorisation date:**

8/11/2024

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### **Manufacturing sites for batch release:**

Ab7 Sante

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### **Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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### **Authorisation number:**

V7006979.00.00

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### **Date of authorisation status change:**

8/11/2024

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### **Reference member state:**

Ireland

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### **Procedure number:**

IE/V/0667/002

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### **Concerned member states:**

France Germany Italy Netherlands Poland Portugal 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

Combined File of all Documents

English (PDF)

Published on: 26/10/2025

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Summary of Product Characteristics