

# Advantage 250 mg Lösung zum Auftropfen für Hunde ( $\geq 10 - < 25$ kg)

Authorised

- Imidacloprid

## Product identification

### **Medicine name:**

Advantage 250 mg Lösung zum Auftropfen für Hunde ( $\geq 10 - < 25$  kg)

ADVANTAGE 250 mg SOLUCION SPOT-ON PARA PERROS ( $\geq 10 - < 25$  KG)

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### **Active substance:**

Imidacloprid

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### **Target species:**

Dog

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### **Route of administration:**

Cutaneous use

## Product details

### **Active substance and strength:**

Imidacloprid

250.00 milligram(s) / 2.50 millilitre(s)

**Pharmaceutical form:**

Cutaneous solution

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

QP53AX17

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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:**

Spain

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**Package description:**

Pack sizes 2.5 ml solution per pipette, Pack containing 1 unit dose pipettes,  
Container: White polypropylene pipettes with caps

Pack sizes 2.5 ml solution per pipette, Pack containing 2 unit dose pipettes,  
Container: White polypropylene pipettes with caps

Pack sizes 2.5 ml solution per pipette, Pack containing 3 unit dose pipettes,  
Container: White polypropylene pipettes with caps

Pack sizes 2.5 ml solution per pipette, Pack containing 4 unit dose pipettes,  
Container: White polypropylene pipettes with caps

Pack sizes 2.5 ml solution per pipette, Pack containing 6 unit dose pipettes,  
Container: White polypropylene pipettes with caps

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Informed consent application (Article 13c of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Elanco Animal Health GmbH

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**Marketing authorisation date:**

2/09/1997

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**Manufacturing sites for batch release:**

KVP Pharma+Veterinaer Produkte GmbH

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

1183 ESP

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**Date of authorisation status change:**

1/01/2023

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**Reference member state:**

Austria

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**Procedure number:**

AT/V/0020/003

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**Concerned member states:**

Denmark Germany Ireland Italy Netherlands Norway Spain Sweden

United Kingdom (Northern Ireland)

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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

Summary of Product Characteristics

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.

## Labelling

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