

# Frento Forte Flohschutztropfen 40 mg Lösung zum Auftropfen für Katzen (<4 kg)

Authorised

- Imidacloprid

## Product identification

**Medicine name:**

Frento Forte Flohschutztropfen 40 mg Lösung zum Auftropfen für Katzen (<4 kg)

---

**Active substance:**

Imidacloprid

---

**Target species:**

Cat

---

**Route of administration:**

Cutaneous use

---

## Product details

**Active substance and strength:**

Imidacloprid

40.00 milligram(s) / 0.40 millilitre(s)

---

**Pharmaceutical form:**

Spot-on solution

---

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

QP53AX17

---

**Legal status of supply:**

Veterinary medicinal product not subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Netherlands

---

**Available in:**

Netherlands

---

**Package description:**

Pack sizes 0.4 ml solution per pipette, Pack containing 1 unit dose pipettes,

Container: White polypropylene pipettes with caps

Pack sizes 0.4 ml solution per pipette, Pack containing 2 unit dose pipettes,

Container: White polypropylene pipettes with caps

Pack sizes 0.4 ml solution per pipette, Pack containing 3 unit dose pipettes,

Container: White polypropylene pipettes with caps

Pack sizes 0.4 ml solution per pipette, Pack containing 4 unit dose pipettes,

Container: White polypropylene pipettes with caps

Pack sizes 0.4 ml solution per pipette, Pack containing 6 unit dose pipettes,

Container: White polypropylene pipettes with caps

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Elanco Animal Health GmbH

---

**Marketing authorisation date:**

7/10/1997

---

**Manufacturing sites for batch release:**

KVP Pharma+Veterinaer Produkte GmbH

---

**Responsible authority:**

Medicines Evaluation Board

---

**Authorisation number:**

REG NL 9214

---

**Date of authorisation status change:**

18/01/2022

---

**Reference member state:**

Austria

---

**Procedure number:**

AT/V/0021/001

---

**Concerned member states:**

Denmark Germany Ireland Italy Netherlands Norway Spain Sweden

United Kingdom (Northern Ireland)

---

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

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

## Summary of Product Characteristics