

# 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)

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**Active substance:**

Imidacloprid

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**Target species:**

Cat

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**Route of administration:**

Cutaneous use

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## Product details

**Active substance and strength:**

Imidacloprid  
40.00 milligram(s) / 0.40 millilitre(s)

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**Pharmaceutical form:**

Spot-on 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:**

Netherlands

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**Available in:**

Netherlands

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Elanco Animal Health GmbH

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**Marketing authorisation date:**

7/10/1997

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**Manufacturing sites for batch release:**

KVP Pharma+Veterinaer Produkte GmbH

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 9214

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**Date of authorisation status change:**

18/01/2022

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**Reference member state:**

Austria

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**Procedure number:**

AT/V/0021/001

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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

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