

# EFFIPRO 402 mg spot-on solution for very large dogs

Authorised

- Fipronil

## Product identification

**Medicine name:**

EFFIPRO 402 mg spot-on solution for very large dogs

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**Active substance:**

Fipronil

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**Target species:**

Dog

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**Route of administration:**

Cutaneous use

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## Product details

**Active substance and strength:**

Fipronil

402.00 milligram(s) / 1.00 Pipette

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**Pharmaceutical form:**

Spot-on solution

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

QP53AX15

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Ireland

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**Package description:**

Box containing 1 pipette of 4.02 mL

Box containing 150 blister packs of 1 pipette of 4.02 mL

Box containing 90 blister packs of 1 pipette of 4.02 mL

Box containing 60 blister packs of 1 pipette of 4.02 mL

Box containing 30 blister packs of 1 pipette of 4.02 mL

Box containing 24 blister packs of 1 pipette of 4.02 mL

Box containing 12 blister packs of 1 pipette of 4.02 mL

Box containing 8 blister packs of 1 pipette of 4.02 mL

Box containing 6 blister packs of 1 pipette of 4.02 mL

Box containing 4 blister packs of 1 pipette of 4.02 mL

Box containing 3 blister packs of 1 pipette of 4.02 mL

Box containing 2 blister packs of 1 pipette of 4.02 mL

Box containing 1 blister pack of 1 pipette of 4.02 mL

Box containing 150 pipettes of 4.02 mL

Box containing 90 pipettes of 4.02 mL

Box containing 60 pipettes of 4.02 mL

Box containing 30 pipettes of 4.02 mL

Box containing 24 pipettes of 4.02 mL

Box containing 12 pipettes of 4.02 mL

Box containing 8 pipettes of 4.02 mL

Box containing 6 pipettes of 4.02 mL

Box containing 4 pipettes of 4.02 mL

Box containing 3 pipettes of 4.02 mL

Box containing 2 pipettes of 4.02 mL

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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:**

Virbac

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**Marketing authorisation date:**

15/05/2009

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**Manufacturing sites for batch release:**

Virbac

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10988/076/004

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**Date of authorisation status change:**

15/05/2009

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**Reference member state:**

France

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**Procedure number:**

FR/V/0377/004

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**Concerned member states:**

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland  
Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg  
Netherlands Poland Portugal Romania Slovakia Slovenia 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)