

# Fyperix 134 mg spot-on solution for dogs

Authorised

- Fipronil

## Product identification

**Medicine name:**

Fyperix 134 mg spot-on solution for dogs

Fyperix vet

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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

134.00 milligram(s) / 0.67 millilitre(s)

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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 not subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Finland

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**Available in:**

Finland

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**Package description:**

(ID6) 40.2 millilitre(s): Box (board) with 30 Beutel (polyethylene terephthalate; aluminium; low-density polyethylene) each with 1 Pipette (polypropylene) with 1.34 millilitre(s)

(ID5) 26.8 millilitre(s): Box (board) with 20 Beutel (polyethylene terephthalate; aluminium; low-density polyethylene) each with 1 Pipette (polypropylene) with 1.34 millilitre(s)

(ID4) 13.4 millilitre(s): Box (board) with 10 Beutel (polyethylene terephthalate; aluminium; low-density polyethylene) each with 1 Pipette (polypropylene) with 1.34 millilitre(s)

(ID3) 8.04 millilitre(s): Box (board) with 6 Beutel (polyethylene terephthalate; aluminium; low-density polyethylene) each with 1 Pipette (polypropylene) with 1.34 millilitre(s)

(ID2) 4.02 millilitre(s): Box (board) with 3 Beutel (polyethylene terephthalate; aluminium; low-density polyethylene) each with 1 Pipette (polypropylene) with 1.34 millilitre(s)

(ID1) 1.34 millilitre(s): Box (board) with Beutel (polyethylene terephthalate; aluminium; low-density polyethylene) with 1 Pipette (polypropylene) with 1.34 millilitre(s)

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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:**

KRKA tovarna zdravil d.d. Novo mesto

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**Marketing authorisation date:**

17/06/2015

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**Manufacturing sites for batch release:**

KRKA tovarna zdravil d.d. Novo mesto

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**Responsible authority:**

Finnish Medicines Agency

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**Authorisation number:**

29827

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**Date of authorisation status change:**

17/06/2015

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0190/003

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**Concerned member states:**

Finland France Italy 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

Summary of Product Characteristics

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