

# Eliminall 134 mg spot-on solution for dogs

Not authorised

- Fipronil

## Product identification

**Medicine name:**

Eliminall 134 mg spot-on solution for 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

134.00 milligram(s) / 1.34 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:**

Surrendered

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**Authorised in:**

Spain

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**Package description:**

(ID6) 1206 millilitre(s): Box (Cardboard) with 30 Beutel (Low Density PolyEthylene; PolyEthylene TerePhthalate; Aluminium) each with 30 Pipette (PolyPropylene) each with 1.34 millilitre(s)

(ID5) 536 millilitre(s): Box (Cardboard) with 20 Beutel (Low Density PolyEthylene; PolyEthylene TerePhthalate; Aluminium) each with 20 Pipette (PolyPropylene) each with 1.34 millilitre(s)

(ID4) 134 millilitre(s): Box (Cardboard) with 10 Beutel (Low Density PolyEthylene; PolyEthylene TerePhthalate; Aluminium) each with 10 Pipette (PolyPropylene) each with 1.34 millilitre(s)

(ID3) 48.24 millilitre(s): Box (Cardboard) with 6 Beutel (Low Density PolyEthylene; PolyEthylene TerePhthalate; Aluminium) each with 6 Pipette (PolyPropylene) each with 1.34 millilitre(s)

(ID2) 12.06 millilitre(s): Box (Cardboard) with 3 Beutel (Low Density PolyEthylene; PolyEthylene TerePhthalate; Aluminium) each with 3 Pipette (PolyPropylene) each with 1.34 millilitre(s)

(ID1) 1.34 millilitre(s): Box (Cardboard) with Beutel (Low Density PolyEthylene; PolyEthylene TerePhthalate; Aluminium) 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:**

14/11/2011

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**Manufacturing sites for batch release:**

KRKA tovarna zdravil d.d. Novo mesto

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

2406 ESP

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**Date of authorisation status change:**

20/01/2025

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0189/003

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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.