

# FYPERIX COMBO 67 MG/60,3 MG SPOT-ON SOLUTION FOR SMALL DOGS

Authorised

- Fipronil
- Methoprene

## Product identification

**Medicine name:**

FYPERIX COMBO 67 MG/60,3 MG SPOT-ON SOLUTION FOR SMALL DOGS

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**Active substance:**

Fipronil

Methoprene

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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

67.00 milligram(s) / 1.00 Pipette

Methoprene  
60.30 milligram(s) / 1.00 Pipette

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**Pharmaceutical form:**

Spot-on solution

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

QP53AX65

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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:**

White polypropylene unit-dose pipette with polyethylene or polyoxymethylene closure with spike packaged into laminated triplex bag composed of polyester, aluminium and polyethylene. Cardboard box containing 1 pipette.

White polypropylene unit-dose pipette with polyethylene or polyoxymethylene closure with spike packaged into laminated triplex bag composed of polyester, aluminium and polyethylene. Cardboard box containing 30 pipettes.

White polypropylene unit-dose pipette with polyethylene or polyoxymethylene closure with spike packaged into laminated triplex bag composed of polyester, aluminium and polyethylene. Cardboard box containing 10 pipettes.

White polypropylene unit-dose pipette with polyethylene or polyoxymethylene closure with spike packaged into laminated triplex bag composed of polyester, aluminium and polyethylene. Cardboard box containing 6 pipettes.

White polypropylene unit-dose pipette with polyethylene or polyoxymethylene closure with spike packaged into laminated triplex bag composed of polyester, aluminium and polyethylene. Cardboard box containing 3 pipettes.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) 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:**

23/05/2016

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**Manufacturing sites for batch release:**

TAD Pharma GmbH

KRKA tovarna zdravil d.d. Novo mesto

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 117510

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**Date of authorisation status change:**

5/04/2022

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**Reference member state:**

France

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**Procedure number:**

FR/V/0333/002

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**Concerned member states:**

Austria Belgium Czechia Estonia Germany Greece Italy Netherlands Poland  
Portugal Romania Spain 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

English (PDF)

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