# Fypryst Combo 268 mg/ 241.2 mg spot-on solution for large dogs

Authorised

- (S)-Methoprene
- Fipronil

## Product identification

#### **Medicine name:**

Fypryst Combo 268 mg/ 241.2 mg spot-on solution for large dogs Фиприст Комбо 268 mg/ 241,2 mg спот-он разтвор за кучета от едри породи

#### **Active substance:**

(S)-Methoprene

Fipronil

## **Target species:**

Dog

#### Route of administration:

Spot-on use

# **Product details**

## **Active substance and strength:**

(S)-Methoprene 241.20 milligram(s) / 2.68 millilitre(s)

**Fipronil** 

268.00 milligram(s) / 2.68 millilitre(s)

#### **Pharmaceutical form:**

Spot-on solution

## Withdrawal period by route of administration:

#### **Spot-on use:**

**.** . . .

# Dog

## Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AX65

### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

Bulgaria

## Package description:

White polypropylene single-dose pipette packaged in aluminium foil sachets.

Cardboard box containing 1pipette.

White polypropylene single-dose pipette packaged in aluminium foil sachets.

Cardboard box containing 10 pipettes

White polypropylene single-dose pipette packaged in aluminium foil sachets.

Cardboard box containing 6 pipettes

White polypropylene single-dose pipette packaged in aluminium foil sachets.

Cardboard box containing 3 pipettes

# Additional information

# **Entitlement type:**

Marketing Authorisation

# Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:	
KRKA tovarna zdravil d.d. Novo mesto	
Marketing authorisation date:	
24/03/2014	
Manufacturing sites for batch release:	
KRKA tovarna zdravil d.d. Novo mesto	
Responsible authority:	
Bulgarian Food Safety Authority	
Bulgarian rood Safety Authority	
Authorisation number:	
0022-2239	
Date of authorisation status change:	
27/02/2019	
27/02/2019	
Reference member state:	
Hungary	
Dura da da marana da marana	
Procedure number:	
HU/V/0133/003/DC	

## **Concerned member states:**

Bulgaria Croatia Estonia Latvia Lithuania Poland Romania Slovenia

To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>

# **Documents**

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.

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