

Fiprotec 134 mg Spot on Solution for Medium Dogs

Authorised

- Fipronil

Product identification

Medicine name:

Fiprotec 134 mg Spot on Solution for Medium Dogs

Active substance:

Fipronil

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Fipronil

134.00 milligram(s) / 1.34 millilitre(s)

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AX15

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Lithuania

Package description:

Blue pipette composed of a heat-formed shell and a film. 6 pipettes are packed in a cardboard box

Blue pipette composed of a heat-formed shell and a film, enclosed in an aluminium blister. 6 pipettes are packed in a cardboard box

Blue pipette composed of a heat-formed shell and a film. 1 pipette is packed in a cardboard box

Blue pipette composed of a heat-formed shell and a film. 2 pipettes are packed in a cardboard box

Blue pipette composed of a heat-formed shell and a film. 3 pipettes are packed in a cardboard box

Blue pipette composed of a heat-formed shell and a film, enclosed in an aluminium blister. 4 pipettes are packed in a cardboard box

Blue pipette composed of a heat-formed shell and a film, enclosed in an aluminium blister. 1 pipette is packed in a cardboard box

Blue pipette composed of a heat-formed shell and a film, enclosed in an aluminium blister. 2 pipettes are packed in a cardboard box

Blue pipette composed of a heat-formed shell and a film, enclosed in an aluminium blister. 3 pipettes are packed in a cardboard box

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Beaphar B.V.

Marketing authorisation date:

25/06/2014

Manufacturing sites for batch release:

Beaphar B.V.

Laboratorios Calier S.A.

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/14/2228/007-011;028-031

Date of authorisation status change:

27/04/2026

Reference member state:

Netherlands

Procedure number:

NL/V/0274/003

Concerned member states:

Bulgaria Cyprus France Germany Greece Hungary Italy Latvia Lithuania
Poland Portugal Romania Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

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