

Fiprotec 50 mg Spot-On Solution for Cats

Authorised

- Fipronil

Product identification

Medicine name:

Fiprotec 50 mg Spot-On Solution for Cats

Active substance:

Fipronil

Target species:

Cat

Route of administration:

Spot-on use

Product details

Active substance and strength:

Fipronil

50.00 milligram(s) / 0.50 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:

Germany

Available in:

Germany

Package description:

Blue pipette composed of a heat-formed shell and a film, enclosed in an aluminium blister. 3 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. 6 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

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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. 6 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:

19/05/2014

Manufacturing sites for batch release:

Beaphar B.V.

Laboratorios Calier S.A.

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

401920.00.00

Date of authorisation status change:

27/09/2019

Reference member state:

Netherlands

Procedure number:

NL/V/0274/002

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

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