Fiprotec 402 mg Spot on Solution for Extra Large Dogs

Authorised

- Fipronil

Product identification

Medicine name:
Fiprotec 402 mg Spot on Solution for Extra Large Dogs
Fiprotec 402 mg šķīdums pilināšanai uz ādas ļoti liela auguma suņiem

Active substance:
Fipronil

Target species:
Dog

Route of administration:
Spot-on use

Product details

Active substance and strength:
Fipronil
402.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Spot-on solution

Withdrawal period by route of administration:

Spot-on use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AX15

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

Package description:

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. 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. 1 pipette is 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
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, enclosed in an aluminium blister. 4 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. 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 BV

Marketing authorisation date:
9/06/2014

Manufacturing sites for batch release:
Beaphar B.V.
Laboratorios Calier S.A.

Responsible authority:
PVD

Authorisation number:
V/DCP/14/0025

Date of authorisation status change:
9/06/2014

Reference member state:
Netherlands

Procedure number:
NL/V/0274/005

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

Labelling

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.

Summary of Product Characteristics

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