Fiprotec 268 mg Spot on Solution for Large Dogs

Authorised

- Fipronil

Product identification

Medicine name:
Fiprotec 268 mg Spot on Solution for Large Dogs
Fiprotec 268 mg šķīdums pilināšanai uz ādas liela auguma suņiem

Active substance:
Fipronil

Target species:
Dog

Route of administration:
Spot-on use

Product details

Active substance and strength:
Fipronil
268.00 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. 3 pipettes are 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. 3 pipettes are 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. 1 pipette is 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

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/14/0024

Date of authorisation status change:
9/06/2014

Reference member state:
Netherlands

Procedure number:
NL/V/0274/004

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

Package Leaflet

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Labelling

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Summary of Product Characteristics

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

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