Perfikan 26.8 mg/240 mg spot-on solution for very small dogs

- Fipronil
- Permethrin

Product identification

Medicine name: Perfikan 26.8 mg/240 mg spot-on solution for very small dogs CANIGUARD DUO

Active substance:

Fipronil Permethrin

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Fipronil 26.84 milligram(s) / 1.00 Pipette Permethrin 239.80 milligram(s) / 1.00 Pipette

Pharmaceutical form:

Spot-on solution

Withdrawal period by route of administration:

Spot-on use:

- . Dog
 - Not applicable. 9999 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AC54

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Package description:

Transparent multi-layer plastic single-dose pipettes containing 0.44 ml obtained by thermoforming a transparent bottom complex (polyacrylonitrile-methacrylate/ or polyethylene-ethylene vinyl alcohol-polyethylene/polypropylene/cyclic olefin copolymer/ polypropylene) and closed by heat sealing with a lid complex (polyacrylonitrile-methacrylate or polyethylene-ethylene vinyl alcohol-polyethylene / aluminium/ polyethylene-terephthalate). Boxes of 6 pipettes.

Transparent multi-layer plastic single-dose pipettes containing 0.44 ml obtained by thermoforming a transparent bottom complex (polyacrylonitrile-methacrylate/ or polyethylene-ethylene vinyl alcohol-polyethylene/polypropylene/cyclic olefin copolymer/ polypropylene) and closed by heat sealing with a lid complex (polyacrylonitrile-methacrylate or polyethylene-ethylene vinyl alcohol-polyethylene / aluminium/ polyethylene-terephthalate). Boxes of 4 pipettes.

Transparent multi-layer plastic single-dose pipettes containing 0.44 ml obtained by thermoforming a transparent bottom complex (polyacrylonitrile-methacrylate/ or polyethylene-ethylene vinyl alcohol-polyethylene/polypropylene/cyclic olefin copolymer/ polypropylene) and closed by heat sealing with a lid complex (polyacrylonitrile-methacrylate or polyethylene-ethylene vinyl alcohol-polyethylene /

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:

Alfamed

Marketing authorisation date:

21/05/2018

Manufacturing sites for batch release:

Virbac France

Responsible authority:

Ministry Of Health

Authorisation number:

104974

Date of authorisation status change:

21/05/2018

Reference member state:

Portugal

Procedure number:

PT/V/0134/001

Concerned member states:

Italy Spain

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

Documents

Summary of Product Characteristics

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Combined File of all Documents

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