

Zeronil Combo 67 mg/60.3 mg Spot-on Solution for Small Dogs

Authorised

- Fipronil
- (S)-Methoprene

Product identification

Medicine name:

Zeronil Combo 67 mg/60.3 mg Spot-on Solution for Small Dogs

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Active substance:

Fipronil

(S)-Methoprene

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Fipronil

67.00 milligram(s) / 1.00 Pipette

(S)-Methoprene
60.30 milligram(s) / 1.00 Pipette

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AX65

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

A white pipette composed of a heat-formed shell of polypropylene/cyclic olefin copolymer/polypropylene layer andpolyethylene/ethylene vinyl alcohol/polyethylene layer.Box with 160 pipettes in individual foil sachets.

A white pipette composed of a heat-formed shell of polypropylene/cyclic olefin copolymer/polypropylene layer andpolyethylene/ethylene vinyl alcohol/polyethylene layer.Box with 150 pipettes in individual foil sachets.

A white pipette composed of a heat-formed shell of polypropylene/cyclic olefin copolymer/polypropylene layer andpolyethylene/ethylene vinyl alcohol/polyethylene layer.Box with 90 pipettes in individual foil sachets.

A white pipette composed of a heat-formed shell of polypropylene/cyclic olefin copolymer/polypropylene layer andpolyethylene/ethylene vinyl alcohol/polyethylene layer.Box with 60 pipettes in individual foil sachets.

A white pipette composed of a heat-formed shell of polypropylene/cyclic olefin copolymer/polypropylene layer andpolyethylene/ethylene vinyl alcohol/polyethylene layer.Box with 30 pipettes in individual foil sachets.

A white pipette composed of a heat-formed shell of polypropylene/cyclic olefin copolymer/polypropylene layer andpolyethylene/ethylene vinyl alcohol/polyethylene layer.Box with 24 pipettes in individual foil sachets.

A white pipette composed of a heat-formed shell of polypropylene/cyclic olefin copolymer/polypropylene layer andpolyethylene/ethylene vinyl alcohol/polyethylene

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:

Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:

20/04/2018

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10987/126/001

Date of authorisation status change:

20/04/2018

Reference member state:

Ireland

Procedure number:

IE/V/0386/002

Concerned member states:

Austria Bulgaria Cyprus Czechia France Greece Hungary Italy Netherlands
Poland Portugal Romania Slovakia Spain Sweden

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

Documents

Combined File of all Documents