

PestiGon Combo 67 mg / 60.3 mg spot-on solution for small dogs

Authorised

- Fipronil
- (S)-Methoprene

Product identification

Medicine name:

PestiGon Combo 67 mg / 60.3 mg spot-on solution for small dogs

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 not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Estonia

Package description:

0.67 ml pipette, moulded from a film composed of 3 layers: a polypropylene/COC/polypropylene, solvent free lacquer laminate and a copolymer of polyethylene/EVOH/polyethylene. The pipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film and presented in an outer box. Boxes of 24 pipettes. Each pipette is individually sealed in a foil sachet.

0.67 ml pipette, moulded from a film composed of 3 layers: a polypropylene/COC/polypropylene, solvent free lacquer laminate and a copolymer of polyethylene/EVOH/polyethylene. The pipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film and presented in an outer box. Boxes of 12 pipettes. Each pipette is individually sealed in a foil sachet.

0.67 ml pipette, moulded from a film composed of 3 layers: a polypropylene/COC/polypropylene, solvent free lacquer laminate and a copolymer of polyethylene/EVOH/polyethylene. The pipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film and presented in an outer box. Boxes of 8 pipettes. Each pipette is individually sealed in a foil sachet.

0.67 ml pipette, moulded from a film composed of 3 layers: a polypropylene/COC/polypropylene, solvent free lacquer laminate and a copolymer of polyethylene/EVOH/polyethylene. The pipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film and presented in an outer box. Boxes of 6 pipettes. Each pipette is individually sealed in a foil sachet.

0.67 ml pipette, moulded from a film composed of 3 layers: a polypropylene/COC/polypropylene, solvent free lacquer laminate and a copolymer of polyethylene/EVOH/polyethylene. The pipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film and presented in an outer box. Boxes of 4 pipettes. Each pipette is individually sealed in a foil sachet.

0.67 ml pipette, moulded from a film composed of 3 layers: a polypropylene/COC/polypropylene, solvent free lacquer laminate and a copolymer of polyethylene/EVOH/polyethylene. The pipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film and presented in an outer box. Boxes of 3 pipettes. Each pipette is individually sealed in a foil sachet.

0.67 ml pipette, moulded from a film composed of 3 layers: a polypropylene/COC/polypropylene, solvent free lacquer laminate and a copolymer of polyethylene/EVOH/polyethylene. The pipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film and presented in an outer box. Boxes of 2 pipettes. Each pipette is individually sealed in a foil sachet.

0.67 ml pipette, moulded from a film composed of 3 layers: a polypropylene/COC/polypropylene, solvent free lacquer laminate and a copolymer of polyethylene/EVOH/polyethylene. The pipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film and presented in an outer box. Boxes of 1 pipette. Each pipette is individually sealed in a foil sachet.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:

28/03/2017

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

State Agency Of Medicines

Authorisation number:

2023

Date of authorisation status change:

28/03/2017

Reference member state:

Ireland

Procedure number:

IE/V/0363/002

Concerned member states:

Bulgaria Estonia France Greece Hungary Italy Latvia Lithuania Portugal
Romania United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

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