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
PestiGon Combo, 67 mg / 60,3 mg täpilahus väikestele koertele

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 laminateand 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.
0.67 ml pipette, moulded from a film composed of 3 layers: a polypropylene/COC/polypropylene, solvent free lacquer laminateand 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 laminateand 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 laminateand 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 laminateand 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 laminateand 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 laminateand 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 laminateand 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.

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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