Combotec 268 mg/241.2 mg Spot-on Solution for Large Dogs

Authorised

- (S)-Methoprene
- Fipronil

Product identification

Medicine name:
Combotec 268 mg/241.2 mg Spot-on Solution for Large Dogs

Active substance:
(S)-Methoprene

Fipronil

Target species:
Dog

Route of administration:
Cutaneous use

Product details

Active substance and strength:
(S)-Methoprene
241.20 milligram(s) / 1.00 millilitre(s)

Fipronil

268.00 milligram(s) / 1.00 millilitre(s)

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:
Netherlands

Available in:
Netherlands

Package description:
A blue pipette is composed of a heat-formed shell and a film. The blue pipette is enclosed in an aluminum blister. 1 pipette are packed in a cardboard box.
A blue pipette is composed of a heat-formed shell and a film. The blue pipette is enclosed in an aluminum blister. 2 pipettes are packed in a cardboard box.
A blue pipette is composed of a heat-formed shell and a film. The blue pipette is enclosed in an aluminum blister. 3 pipettes are packed in a cardboard box.
A blue pipette is composed of a heat-formed shell and a film. The blue pipette is enclosed in an aluminum blister. 4 pipettes are packed in a cardboard box.
TO BE DELETED - A blue pipette is composed of a heat-formed shell and a film. The blue pipette is enclosed in an aluminum blister. 5 pipettes are packed in a cardboard box.
A blue pipette is composed of a heat-formed shell and a film. The blue pipette is enclosed in an aluminum blister. 6 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 B.V.

Marketing authorisation date:
19/12/2019

Manufacturing sites for batch release:
Beaphar B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 123868

Date of authorisation status change:
19/01/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0251/004

Concerned member states:
Czechia Hungary Italy Malta

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

Documents

Combined File of all Documents

English (PDF)
Published on: 5/02/2026
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