

Eliminall 67 mg spot-on solution for dogs

Authorised

- Fipronil

Product identification

Medicine name:

Eliminall 67 mg spot-on solution for dogs

Active substance:

Fipronil

Target species:

Dog

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Fipronil

67.00 milligram(s) / 0.67 millilitre(s)

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AX15

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Available in:

Greece

Package description:

(ID1) 0.67 millilitre(s): Box (cardboard) with Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) with 1 Pipette (polypropylene) with 0.67 millilitre(s)

(ID2) 6.03 millilitre(s): Box (cardboard) with 3 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 3 Pipette (polypropylene) each with 0.67 millilitre(s)

(ID3) 24.12 millilitre(s): Box (cardboard) with 6 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 6 Pipette (polypropylene) each with 0.67 millilitre(s)

(ID4) 67 millilitre(s): Box (cardboard) with 10 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 10 Pipette (polypropylene) each with 0.67 millilitre(s)

(ID5) 268 millilitre(s): Box (cardboard) with 20 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 20 Pipette (polypropylene) each with 0.67 millilitre(s)

(ID6) 603 millilitre(s): Box (cardboard) with 30 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 30 Pipette (polypropylene) each with 0.67 millilitre(s)

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:

KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:

10/06/2012

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

National Organization For Medicines

Authorisation number:

71095/10-08-2017/K-0190302

Date of authorisation status change:

9/08/2017

Reference member state:

Germany

Procedure number:

DE/V/0189/002

Concerned member states:

Finland Greece Portugal Spain

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

Documents

Combined File of all Documents

2401446-paren-20210316.pdf