

Eliminall 402 mg spot-on solution for dogs

Authorised

- Fipronil

Product identification

Medicine name:

Eliminall 402 mg spot-on solution for dogs

Active substance:

Fipronil

Target species:

Dog

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Fipronil

402.00 milligram(s) / 4.02 millilitre(s)

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AX15

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Finland

Package description:

(ID1) 4.02 millilitre(s): Box (cardboard) with Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) with 1 Pipette (polypropylene) with 4.02 millilitre(s)

(ID2) 36.18 millilitre(s): Box (cardboard) with 3 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 3 Pipette (polypropylene) each with 4.02 millilitre(s)

(ID3) 144.72 millilitre(s): Box (cardboard) with 6 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 6 Pipette (polypropylene) each with 4.02 millilitre(s)

(ID4) 402 millilitre(s): Box (cardboard) with 10 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 10 Pipette (polypropylene) each with 4.02 millilitre(s)

(ID5) 1608 millilitre(s): Box (cardboard) with 20 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 20 Pipette (polypropylene) each with 4.02 millilitre(s)

(ID6) 3618 millilitre(s): Box (cardboard) with 30 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 30 Pipette (polypropylene) each with 4.02 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:

8/05/2012

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Finnish Medicines Agency

Authorisation number:

28890

Date of authorisation status change:

8/05/2012

Reference member state:

Germany

Procedure number:

DE/V/0189/005

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

2401449-paren-20210316.pdf