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 subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

Available in:
Greece

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:
10/06/2012

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
National Organization For Medicines

Authorisation number:
71098/10-08-2017/K-0190305

Date of authorisation status change:
9/08/2017

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