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Eliminall 50 mg spot-on solution for cats

Authorised

- Fipronil

Product identification

Medicine name:
Eliminall 50 mg spot-on solution for cats

Active substance:
Fipronil

Target species:
Cat

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Fipronil
50.00 milligram(s) / 0.50 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) 0.5 millilitre(s): Box (cardboard) with Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) with 1 Pipette (polypropylene) with 0.5 millilitre(s)
(ID2) 4.5 millilitre(s): Box (cardboard) with 3 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 3 Pipette (polypropylene) each with 0.5 millilitre(s)
(ID3) 18 millilitre(s): Box (cardboard) with 6 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 6 Pipette (polypropylene) each with 0.5 millilitre(s)
(ID4) 50 millilitre(s): Box (cardboard) with 10 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 10 Pipette (polypropylene) each with 0.5 millilitre(s)
(ID5) 200 millilitre(s): Box (cardboard) with 20 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 20 Pipette (polypropylene) each with 0.5 millilitre(s)
(ID6) 450 millilitre(s): Box (cardboard) with 30 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 30 Pipette (polypropylene) each with 0.5 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:
28886

Date of authorisation status change:
8/05/2012

Reference member state:
Germany

Procedure number:
DE/V/0189/001

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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

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