Eliminall 50 mg spot-on solution for cats

Not 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:
Surrendered

Authorised in:
Spain

Package description:
(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)
(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)
(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)
(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)
(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)
(ID1) 0.5 millilitre(s): Box (Cardboard) with Beutel (Low Density PolyEthylene; PolyEthylene TerePhthalate; Aluminium) with 1 Pipette (PolyPropylene) 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:
14/11/2011

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

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
2404 ESP

Date of authorisation status change:
20/01/2025

Reference member state:
Germany

Procedure number:
DE/V/0189/001

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

Documents

Labelling

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

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.