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 not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Portugal

Available in:
Portugal

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:
18/10/2011

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

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
376/02/11DFVPT

Date of authorisation status change:
6/02/2026

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

English (PDF)
Published on: 16/04/2026
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