

Eliminall 134 mg spot-on solution for dogs

Authorised

- Fipronil

Product identification

Medicine name:

Eliminall 134 mg spot-on solution for dogs

Active substance:

Fipronil

Target species:

Dog

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Fipronil

134.00 milligram(s) / 1.34 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:

Germany

Package description:

(ID1) 1.34 millilitre(s): Box (cardboard) with Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) with 1 Pipette (polypropylene) with 1.34 millilitre(s)

(ID2) 12.06 millilitre(s): Box (cardboard) with 3 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 3 Pipette (polypropylene) each with 1.34 millilitre(s)

(ID3) 48.24 millilitre(s): Box (cardboard) with 6 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 6 Pipette (polypropylene) each with 1.34 millilitre(s)

(ID4) 134 millilitre(s): Box (cardboard) with 10 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 10 Pipette (polypropylene) each with 1.34 millilitre(s)

(ID5) 536 millilitre(s): Box (cardboard) with 20 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 20 Pipette (polypropylene) each with 1.34 millilitre(s)

(ID6) 1206 millilitre(s): Box (cardboard) with 30 Beutel (low-density polyethylene; polyethylene terephthalate; aluminium) each with 30 Pipette (polypropylene) each with 1.34 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:

4/11/2011

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

401446.01.00

Date of authorisation status change:

5/06/2017

Reference member state:

Germany

Procedure number:

DE/V/0189/003

Concerned member states:

Finland Greece Portugal Spain

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Documents

Combined File of all Documents

English (PDF)

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