Eliminall 402 mg spot-on solution for dogs

Not authorised

- Fipronil

Product identification

Medicine name:
Eliminall 402 mg spot-on solution for dogs
ELIMINALL 402 mg SOLUCION PARA UNCION DORSAL PUNTUAL PARA PERROS

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

Withdrawal period by route of administration:

Cutaneous use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AX15

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

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Authorisation status:
Surrendered

Authorised in:
Spain

Package description:
(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)
(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)
(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)
(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)
(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)
(ID1) 4.02 millilitre(s): Box (Cardboard) with Beutel (Low Density PolyEthylene; PolyEthylene TerePhthalate; Aluminium) with 1 Pipette (PolyPropylene) 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:
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:
2408 ESP

Date of authorisation status change:
20/01/2025

Reference member state:
Germany

Procedure number:
DE/V/0189/005

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

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