Prinocate 40 mg/4 mg spot-on solution for small cats and ferrets

Authorised

- Moxidectin
- Imidacloprid

Product identification

Medicine name:
Prinocate 40 mg/4 mg spot-on solution for small cats and ferrets
Prinocate 40 mg / 0,4 ml + 4 mg / 0,4 ml Roztwór do nakrapiania

Active substance:
Moxidectin
Imidacloprid

Target species:
Cat
Ferret

Route of administration:
Spot-on use

Product details

Active substance and strength:
Moxidectin
4.00 milligram(s) / 1.00 Pipette

Imidacloprid
40.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:
Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AB52

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Poland

Available in:
Poland

Package description:
A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) orpolyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyetylene (LDPE).Cardboard box containing 48 pipettes.
A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) orpolyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyetylene (LDPE).Cardboard box containing 24 pipettes.
A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) orpolyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyetylene (LDPE).Cardboard box containing 6 pipettes.
A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) orpolyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyetylene (LDPE).Cardboard box containing 4 pipettes.

A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) orpolyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyetylene (LDPE).Cardboard box containing 3 pipettes.
A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) orpolyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium(Al) and low density polyetylene (LDPE).Cardboard box containing 1 pipette.

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:
27/01/2020

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

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2939

Date of authorisation status change:
27/01/2020

Reference member state:
Ireland

Procedure number:

IE/V/0392/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia France Greece Hungary Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 16/01/2026
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Labelling

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Package Leaflet

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

Combined File of all Documents

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