Prinocate 400 mg/100 mg spot-on solution for extra-large dogs

Authorised

- Moxidectin
- Imidacloprid

Product identification

Medicine name:
Prinocate 400 mg/100 mg spot-on solution for extra-large dogs
Prinocate 400 mg/100 mg solutie spot-on pentru caini de talie foarte mare

Active substance:
Moxidectin
Imidacloprid

Target species:
Dog

Route of administration:
Spot-on use

Product details

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

Imidacloprid
400.00 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:
Romania

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 1 pipette.
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 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 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 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 48 pipettes.

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:
28/01/2020

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

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
250022

Date of authorisation status change:
25/03/2025

Reference member state:
Ireland

Procedure number:
IE/V/0392/006

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

Combined File of all Documents

English (PDF)
Published on: 23/02/2025
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