Prinocate 80 mg/8 mg spot-on solution for large cats

Authorised

- Imidacloprid
- Moxidectin

Product identification

Medicine name:
Prinocate 80 mg/8 mg spot-on solution for large cats

Active substance:
Imidacloprid

Moxidectin

Target species:
Cat

Route of administration:
Spot-on use

Product details

Active substance and strength:
Imidacloprid
80.00 milligram(s) / 1.00 Pipette

Moxidectin

8.00 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:
Netherlands

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:
6/02/2020

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

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 124515

Date of authorisation status change:
26/01/2022

Reference member state:
Ireland

Procedure number:
IE/V/0392/002

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

Combined File of all Documents

English (PDF)
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Summary of Product Characteristics

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