

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

Authorised

- Moxidectin
- Imidacloprid

Product identification

Medicine name:

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

Prinocate 80 mg/8 mg διάλυμα για επίχυση σε σημείο για μεγαλόσωμες γάτες

Active substance:

Moxidectin

Imidacloprid

Target species:

Cat

Route of administration:

Spot-on use

Product details

Active substance and strength:

Moxidectin

8.00 milligram(s) / 1.00 Pipette

Imidacloprid
80.00 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:

Cyprus

Package description:

A white polypropylene (PP) unit dose pipette with a closure with a spike composed of high density polyethylene (HDPE) or polyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium (Al) and low density polyethylene (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) or polyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium (Al) and low density polyethylene (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) or polyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium (Al) and low density polyethylene (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) or polyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium (Al) and low density polyethylene (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) or polyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium (Al) and low density polyethylene (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) or polyoxymethylene (POM) or polypropylene (PP) packed into a laminated triplex bag composed of polyester (PETP), aluminium (Al) and low density polyethylene (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:

15/01/2020

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

CY00772V

Date of authorisation status change:

15/01/2020

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

Summary of Product Characteristics

Combined File of all Documents

English (PDF)

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