

Prazpronto 40 mg Spot-on solution for medium cats

Not
authorised

- Praziquantel

Product identification

Medicine name:

Prazpronto 40 mg Spot-on solution for medium cats

Active substance:

Praziquantel

Target species:

Cat

Route of administration:

Spot-on use

Product details

Active substance and strength:

Praziquantel

40.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AA01

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Spain

Package description:

A white pipette composed of a heat-formed shell composed of polypropylene/cyclic olefin copolymer/ethylene vinyl alcohol/polypropylene layer. Carton containing 1 pipette in individual foil sachets.

A white pipette composed of a heat-formed shell composed of polypropylene/cyclic olefin copolymer/ethylene vinyl alcohol/polypropylene layer. Carton containing 2 pipettes in individual foil sachets.

A white pipette composed of a heat-formed shell composed of polypropylene/cyclic olefin copolymer/ethylene vinyl alcohol/polypropylene layer. Carton containing 3 pipettes in individual foil sachets.

A white pipette composed of a heat-formed shell composed of polypropylene/cyclic olefin copolymer/ethylene vinyl alcohol/polypropylene layer. Carton containing 4 pipettes in individual foil sachets.

A white pipette composed of a heat-formed shell composed of polypropylene/cyclic olefin copolymer/ethylene vinyl alcohol/polypropylene layer. Carton containing 6 pipettes in individual foil sachets.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:

26/04/2021

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

Spanish Agency For Medicines And Health Products

Authorisation number:

4000 ESP

Date of authorisation status change:

18/05/2023

Reference member state:

Ireland

Procedure number:

IE/V/0633/002

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 3/05/2024

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Package Leaflet

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Labelling

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