

# 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

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**Active substance:**

Praziquantel

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**Target species:**

Cat

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**Route of administration:**

Spot-on use

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## Product details

**Active substance and strength:**

Praziquantel  
40.00 milligram(s) / 1.00 Pipette

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**Pharmaceutical form:**

Spot-on solution

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QP52AA01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

Germany

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Chanelle Pharmaceuticals Manufacturing Limited

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**Marketing authorisation date:**

5/10/2020

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**Manufacturing sites for batch release:**

Chanelle Pharmaceuticals Manufacturing Limited

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

402725.00.00

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**Date of authorisation status change:**

15/05/2023

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0633/002

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics