

Prazpronto 60 mg Spot-on solution for large cats

Not
authorised

- Praziquantel

Product identification

Medicine name:

Prazpronto 60 mg Spot-on solution for large cats

PRAZPRONTO 60 mg SOLUCION SPOT-ON PARA GATOS GRANDES

Active substance:

Praziquantel

Target species:

Cat

Route of administration:

Spot-on use

Product details

Active substance and strength:

Praziquantel

60.00 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:

4001 ESP

Date of authorisation status change:

18/05/2023

Reference member state:

Ireland

Procedure number:

IE/V/0633/003

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

[Download](#)

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.