Prazpronto 20mg Spot-on solution for small Cats and Kittens

Not authorised

- Praziquantel

Product identification

Medicine name:
Prazpronto 20mg Spot-on solution for small Cats and Kittens
PRAZPRONTO

Active substance:
Praziquantel

Target species:
Cat

Route of administration:
Spot-on use

Product details

Active substance and strength:
Praziquantel
20.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:
Spot-on solution

Withdrawal period by route of administration:

Spot-on use:

- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AA01

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

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Authorisation status:

Revoked

Authorised in:

Italy

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:
1/01/2022

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Ltd

Responsible authority:
Ministry Of Health

Authorisation number:
105347

Date of authorisation status change:
11/05/2023

Reference member state:
Ireland

Procedure number:
IE/V/0633/001

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

Documents

Summary of Product Characteristics