Dehinel 230 mg/20 mg film-coated tablets for cats

Authorised

- Praziquantel
- Pyrantel embonate

Product identification

Medicine name:
Dehinel 230 mg/20 mg film-coated tablets for cats
ANTHELMIN 230 mg/20 mg επικαλυμμένα με λεπτό υμένιο δισκία για γάτες

Active substance:
Praziquantel

Pyrantel embonate

Target species:
Cat

Route of administration:
Oral use

Product details

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

Pyrantel embonate
230.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Film-coated tablet

Withdrawal period by route of administration:
Oral use:

- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AA51

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

Package description:
(ID3) 10 Film-coated tablet: unspecified outer container with 1 Blister (Orientated PolyAmide; Aluminium; PolyVinyl Chloride) with 10 Film-coated tablet, closed with Foil (Aluminium)
(ID2) 4 Film-coated tablet: unspecified outer container with 2 Blister (Orientated PolyAmide; Aluminium; PolyVinyl Chloride) each with 2 Film-coated tablet, closed with Foil (Aluminium)
(ID1) 2 Film-coated tablet: unspecified outer container with 1 Blister (Orientated PolyAmide; Aluminium; PolyVinyl Chloride) with 2 Film-coated tablet, closed with Foil (Aluminium)

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:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
9/07/2018

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto
TAD Pharma GmbH
Krka-Farma d.o.o.
Krka-Farma d.o.o.

Responsible authority:
National Organization For Medicines

Authorisation number:
71598/04-07-2023/K-0227401

Date of authorisation status change:
3/07/2023

Reference member state:
Germany

Procedure number:
DE/V/0160/002

Concerned member states:
Cyprus France Greece Italy Poland

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

Documents

Combined File of all Documents

Summary of Product Characteristics

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

Package Leaflet

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