

# 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

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### **Active substance:**

Praziquantel

Pyrantel embonate

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### **Target species:**

Cat

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### **Route of administration:**

Oral use

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## Product details

### **Active substance and strength:**

Praziquantel

20.00 milligram(s) / 1.00 Tablet

Pyrantel embonate  
230.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Film-coated tablet

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

QP52AA51

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Cyprus

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**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)

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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:**

KRKA tovarna zdravil d.d. Novo mesto

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**Marketing authorisation date:**

19/08/2018

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**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.

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**Responsible authority:**

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

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**Authorisation number:**

CY00657V

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**Date of authorisation status change:**

19/08/2018

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0160/002

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**Concerned member states:**

Cyprus France Greece Italy Poland

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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

Combined File of all Documents

English (PDF)

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