

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AA51

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Cyprus

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:

19/08/2018

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:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

CY00657V

Date of authorisation status change:

19/08/2018

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

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Documents

Combined File of all Documents

English (PDF)

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