

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

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 not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

(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)

(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)

(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)

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:

31/10/2017

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:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/0319717 6/2017

Date of authorisation status change:

6/12/2022

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

Summary of Product Characteristics

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

Package Leaflet and Labelling

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

Combined File of all Documents

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