

CANIHELMIN PLUS

Authorised

- Praziquantel
- Pyrantel embonate
- Febantel

Product identification

Medicine name:

CANIHELMIN PLUS

Active substance:

Praziquantel
Pyrantel embonate
Febantel

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Praziquantel
50.00 milligram(s) / 1.00 Tablet
Pyrantel embonate

144.00 milligram(s) / 1.00 Tablet

Febantel

150.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AA51

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Package description:

Available only in Spanish

Available only in Spanish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Genera d.d.

Marketing authorisation date:

23/06/2016

Manufacturing sites for batch release:

Genera d.d.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

3423 ESP

Date of authorisation status change:

1/01/2023

Reference member state:

Spain

Procedure number:

ES/V/0245/001

Concerned member states:

Bulgaria Czechia Romania

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 22/12/2023

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Package Leaflet

English (PDF)

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Labelling

English (PDF)

Published on: 22/12/2023

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Combined File of all Documents

eu-PUAR-esv0245001-dcp-canihelmin-plus-en.pdf

es-puar-canihelmin-plus-es.pdf