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ZEROCIT 50 mg TABLETS FOR DOGS

Authorised

- Praziquantel

Product identification

Medicine name:
ZEROCIT 50 mg TABLETS FOR DOGS

Active substance:
Praziquantel

Target species:
Dog

Route of administration:
Oral use

Product details

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

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AA01

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Carton containing 2 blisters of 10 tablets (20 tablets)
Carton containing 4 blisters of 10 tablets (40 tablets)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Ashish Life Science Holding B.V.

Marketing authorisation date:
13/05/2025

Manufacturing sites for batch release:
Pantex Holland B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 132317

Date of authorisation status change:

13/03/2025

Reference member state:
Spain

Procedure number:
ES/V/0437/001

Concerned member states:
Hungary Ireland Italy Netherlands

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

Documents

Combined File of all Documents

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Summary of Product Characteristics

Package Leaflet

Labelling