

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:

Ireland

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:

11/10/2024

Manufacturing sites for batch release:

Pantex Holland B.V.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA25433/001/001

Date of authorisation status change:

11/10/2024

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

Summary of Product Characteristics

Package Leaflet

Labelling