

Quanifen (50 mg Praziquantel / 500 mg Fenbendazole) Tablets for Cats and Dogs

Not
authorised

- Praziquantel
- Fenbendazole

Product identification

Medicine name:

Quanifen (50 mg Praziquantel / 500 mg Fenbendazole) Tablets for Cats and Dogs

Active substance:

Praziquantel
Fenbendazole

Target species:

Dog
Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Praziquantel

50.00 milligram(s) / 1.00 Tablet

Fenbendazole

500.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AA51

Legal status of supply:

Medicinal product on medical prescription for renewable delivery

Authorisation status:

Revoked

Authorised in:

Austria

Package description:

Strips: 30 µ aluminium foil coated with 35 gsm extruded polythene.Blisters: Foil blisters (aluminium/aluminium)Pack sizes:Strips and blisters: 96 tablets

Strips: 30 µ aluminium foil coated with 35 gsm extruded polythene.Blisters: Foil blisters (aluminium/aluminium)Pack sizes:Strips and blisters: 120 tablets

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White high density polyethylene (HDPE) containers with a white polypropylene child resistant tamper evident cap.Pack sizes:Containers: 20 tablets

Strips: 30 µ aluminium foil coated with 35 gsm extruded polythene.Blisters: Foil blisters (aluminium/aluminium)Pack sizes:Strips and blisters: 24 tablets

White high density polyethylene (HDPE) containers with a white polypropylene child resistant tamper evident cap.Pack sizes:Containers: 30 tablets

White high density polyethylene (HDPE) containers with a white polypropylene child resistant tamper evident cap.Pack sizes:Containers: 60 tablets

White high density polyethylene (HDPE) containers with a white polypropylene child resistant tamper evident cap.Pack sizes:Containers: 100 tablets

Strips: 30 µ aluminium foil coated with 35 gsm extruded polythene.Blisters: Foil blisters (aluminium/aluminium)Pack sizes:Strips and blisters: 50 tablets

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:

23/11/2005

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

8-00659

Date of authorisation status change:

9/02/2011

Reference member state:

Ireland

Procedure number:

IE/V/0175/001

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 13/07/2025

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Labelling

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Package Leaflet

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Combined File of all Documents