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

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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)

Labelling

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

Package Leaflet

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

Combined File of all Documents