Drontal Plus Tablets

Authorised

- Praziquantel
- Pyrantel embonate
- Febantel

Product identification

Medicine name:
Drontal Plus Tablets

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 not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Ireland

Package description:
Container: Aluminium foil blister or polyethylene-coated aluminium blister.Container colour: Silver or white colouredContainer sizes: Cartons containing 100 tablets.Contents: Pale yellow tablets
Container: Aluminium foil blister or polyethylene-coated aluminium blister.Container colour: Silver or white colouredContainer sizes: Cartons containing 20 tablets.Contents: Pale yellow tablets
Container: Aluminium foil blister or polyethylene-coated aluminium blister.Container colour: Silver or white colouredContainer sizes: Cartons containing 6 tablets.Contents: Pale yellow tablets
Container: Aluminium foil blister or polyethylene-coated aluminium blister.Container colour: Silver or white colouredContainer sizes: Cartons containing 4 tablets.Contents: Pale yellow tablets
Container: Aluminium foil blister or polyethylene-coated aluminium blister.Container colour: Silver or white colouredContainer sizes: Cartons containing 2 tablets.Contents: Pale yellow tablets

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Vetoquinol S.A.

Marketing authorisation date:
12/10/1990

Manufacturing sites for batch release:
KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10521/006/001

Date of authorisation status change:
12/10/1990

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

Documents

Summary of Product Characteristics