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Drontal Cat film-coated tablets

Authorised

- Praziquantel
- Pyrantel embonate

Product identification

Medicine name:
Drontal Cat film-coated tablets

Active substance:
Praziquantel

Pyrantel embonate

Target species:
Cat

Route of administration:
Oral use

Product details

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

Pyrantel embonate
230.00 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

Available in:
Ireland

Package description:
HISTORICAL PACK Container material: Aluminium foil blisterClosure: Heat sealContainer colour: Silver or white colouredContainer sizes: Cartons containing 2 tablets
HISTORICAL PACK Container material: Aluminium foil blisterClosure: Heat sealContainer colour: Silver or white colouredContainer sizes: Cartons containing 24 tablets
HISTORICAL PACK Container material: Aluminium foil blisterClosure: Heat sealContainer colour: Silver or white colouredContainer sizes: Cartons containing 96 tablets
Container material: Polyethylene-coated aluminium blisterClosure: Heat sealContainer colour: Silver or white colouredContainer sizes: Cartons containing 2 tablets
Container material: Polyethylene-coated aluminium blisterClosure: Heat sealContainer colour: Silver or white colouredContainer sizes: Cartons containing 24 tablets
Container material: Polyethylene-coated aluminium blister Closure: Heat sealContainer colour: Silver or white colouredContainer sizes: Cartons containing 96 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:
22/09/1995

Manufacturing sites for batch release:
KVP Pharma+Veterinaer Produkte GmbH
Vetoquinol S.A.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10521/007/001

Date of authorisation status change:
22/09/1995

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

Documents

Summary of Product Characteristics