Metrocare 250 mg tablets for dogs and cats

Authorised

- Metronidazole

Product identification

Medicine name:
Metrocare 250 mg tablets for dogs and cats

Active substance:
Metronidazole

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Metronidazole
250.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01XD01
QP51AA01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
PVC Aluminium Oriented polyamide Aluminium blister packs.Cardboard box of 7 blisters. Pack size 70 tablets.
PVC Aluminium Oriented polyamide Aluminium blister packs. Cardboard box of 9 blisters. Pack size 90 tablets.
PVC Aluminium Oriented polyamide Aluminium blister packs. Cardboard box of 8 blisters. Pack size 80 tablets.
PVC Aluminium Oriented polyamide Aluminium blister packs. Cardboard box of 6 blisters. Pack size 60 tablets.
PVC Aluminium Oriented polyamide Aluminium blister packs. Cardboard box of 50 blisters. Pack size 500 tablets.
PVC Aluminium Oriented polyamide Aluminium blister packs. Cardboard box of 5 blisters. Pack size 50 tablets.
PVC Aluminium Oriented polyamide Aluminium blister packs. Cardboard box of 4 blisters. Pack size 40 tablets.
PVC Aluminium Oriented polyamide Aluminium blister packs. Cardboard box of 3 blisters. Pack size 30 tablets.
PVC Aluminium Oriented polyamide Aluminium blister packs. Cardboard box of 25 blisters. Pack size 250 tablets.
PVC Aluminium Oriented polyamide Aluminium blister packs. Cardboard box of 2 blisters. Pack size 20 tablets.
PVC Aluminium Oriented polyamide Aluminium blister packs. Cardboard box of 10 blisters. Packsize 100 tablets.

PVC Aluminium Oriented polyamide Aluminium blister packs. Cardboard box of 1 blister. Pack size 10 tablets.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ecuphar

Marketing authorisation date:
13/09/2019

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10491/012/001

Date of authorisation status change:
13/09/2019

Reference member state:
Netherlands

Procedure number:
NL/V/0239/001

Concerned member states:
Austria Belgium Czechia Denmark Estonia Finland France Germany Hungary Ireland Luxembourg Norway Poland Portugal Romania Slovakia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents