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:
Germany

Available in:
Germany

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:
17/07/2019

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402554.00.00

Date of authorisation status change:
17/07/2019

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)

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Documents

Combined File of all Documents

English (PDF)
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