

# Metrocare 250 mg tablets for dogs and cats

Authorised

- Metronidazole

## Product identification

**Medicine name:**

Metrocare 250 mg tablets for dogs and cats

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**Active substance:**

Metronidazole

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**Target species:**

Dog

Cat

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Metronidazole

250.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01XD01

QP51AA01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Poland

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**Available in:**

Poland

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ecuphar

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**Marketing authorisation date:**

16/07/2020

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**Manufacturing sites for batch release:**

Lelypharma B.V.

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

2995

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**Date of authorisation status change:**

16/07/2020

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0239/001

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**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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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### 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.

### Summary of Product Characteristics

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

### Combined File of all Documents