

# Metrocare 500 mg tablets for dogs and cats

Authorised

- Metronidazole

## Product identification

**Medicine name:**

Metrocare 500 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

500.00 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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

QJ01XD01

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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:**

Netherlands

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**Available in:**

Netherlands

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**Package description:**

10 tablets - 1 PVC-Aluminium-Oriented polyamide -Aluminium blister pack in a cardboard box.

90 tablets - 9 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.

250 tablets - 25 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.

30 tablets - 3 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.

100 tablets - 10 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.

20 tablets - 2 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.

70 tablets - 7 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.

80 tablets - 8 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.

40 tablets - 4 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.

60 tablets - 6 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.

50 tablets - 5 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.

500 tablets - 50 PVC-Aluminium-Oriented polyamide -Aluminium blisters pack in a cardboard box.

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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:**

19/07/2019

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**Manufacturing sites for batch release:**

Lelypharma B.V.

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 122686

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**Date of authorisation status change:**

26/01/2022

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0239/002

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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

Combined File of all Documents

English (PDF)

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Final PuAR Metrocare 250 mg 500 mg REG NL 122684 122686.pdf