

# Metrobactin 500 mg tablets for cats and dogs

Authorised

- Metronidazole

## Product identification

**Medicine name:**

Metrobactin 500 mg tablets for cats and dogs

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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:**

Portugal

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**Available in:**

Portugal

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**Package description:**

Aluminium - PVC/PE/PVDC blister. Cardboard box containing 10 boxes, each containing 10 blisters of 10 tablets.

Aluminium - PVC/PE/PVDC blister. Cardboard box containing 10 boxes, each containing 1 blister of 10 tablets.

Aluminium - PVC/PE/PVDC blister. Cardboard box of 5 blisters of 10 tablets

Aluminium - PVC/PE/PVDC blister. Cardboard box of 7 blisters of 10 tablets

Aluminium - PVC/PE/PVDC blister. Cardboard box of 6 blisters of 10 tablets

Aluminium - PVC/PE/PVDC blister. Cardboard box of 50 blisters of 10 tablets

Aluminium - PVC/PE/PVDC blister. Cardboard box of 1 blister of 10 tablets

Aluminium - PVC/PE/PVDC blister. Cardboard box of 9 blisters of 10 tablets

Aluminium - PVC/PE/PVDC blister. Cardboard box of 8 blisters of 10 tablets

Aluminium - PVC/PE/PVDC blister. Cardboard box of 10 blisters of 10 tablets

Aluminium - PVC/PE/PVDC blister. Cardboard box of 4 blisters of 10 tablets

Aluminium - PVC/PE/PVDC blister. Cardboard box of 3 blisters of 10 tablets

Aluminium - PVC/PE/PVDC blister. Cardboard box of 25 blisters of 10 tablets

Aluminium - PVC/PE/PVDC blister. Cardboard box of 2 blister of 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:**

Dechra Regulatory B.V.

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**Marketing authorisation date:**

27/10/2015

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**Manufacturing sites for batch release:**

Genera d.d.

Lelypharma B.V.

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**Responsible authority:**

Directorate General For Food And Veterinary

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**Authorisation number:**

956/02/15DFVPT

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**Date of authorisation status change:**

24/01/2025

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0193/002

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania  
Luxembourg Norway Poland Portugal Romania Slovakia Slovenia 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

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