

# FUNGICONAZOL 400 MG TABLETS FOR DOGS

Authorised

- Ketoconazole

## Product identification

**Medicine name:**

FUNGICONAZOL 400 MG TABLETS FOR DOGS

FUNGICONAZOL 400

**Active substance:**

Ketoconazole

**Target species:**

Dog

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Ketoconazole

400.00 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**

Tablet

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

QJ02AB02

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Italy

---

**Package description:**

Cardboard box of 1 Aluminium/PVC/PE/PVDC blister containing 10 tablets each.  
Cardboard box of 10 Aluminium/PVC/PE/PVDC blisters containing 10 tablets each.  
Cardboard box of 9 Aluminium/PVC/PE/PVDC blisters containing 10 tablets each.  
Cardboard box of 8 Aluminium/PVC/PE/PVDC blisters containing 10 tablets each.  
Cardboard box of 7 Aluminium/PVC/PE/PVDC blisters containing 10 tablets each.  
Cardboard box of 6 Aluminium/PVC/PE/PVDC blisters containing 10 tablets each.  
Cardboard box of 5 Aluminium/PVC/PE/PVDC blisters containing 10 tablets each.  
Cardboard box of 4 Aluminium/PVC/PE/PVDC blisters containing 10 tablets each.  
Cardboard box of 3 Aluminium/PVC/PE/PVDC blisters containing 10 tablets each.  
Cardboard box of 2 Aluminium/PVC/PE/PVDC blisters containing 10 tablets each.

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Dechra Regulatory B.V.

---

**Marketing authorisation date:**

18/09/2015

---

**Manufacturing sites for batch release:**

Genera d.d.  
Lelypharma B.V.

---

**Responsible authority:**

Ministry Of Health

---

**Authorisation number:**

104686

---

**Date of authorisation status change:**

18/09/2015

---

**Reference member state:**

France

---

**Procedure number:**

FR/V/0263/002

---

**Concerned member states:**

Austria Belgium Croatia Czechia Denmark Estonia Finland Greece Hungary  
Iceland Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway  
Poland Portugal Romania Slovakia Slovenia Spain Sweden  
United Kingdom (Northern Ireland)

---

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

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