

FUNGICONAZOL 200 MG TABLETS FOR DOGS

Authorised

- Ketoconazole

Product identification

Medicine name:

FUNGICONAZOL 200 MG TABLETS FOR DOGS

Fungiconazol 200 mg Tabletten für Hunde

Active substance:

Ketoconazole

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Ketoconazole

200.00 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:

Austria

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:

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

Marketing authorisation holder:

Dechra Regulatory B.V.

Marketing authorisation date:

3/10/2014

Manufacturing sites for batch release:

Genera d.d.
Lelypharma B.V.

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

835813

Date of authorisation status change:

3/10/2014

Reference member state:

France

Procedure number:

FR/V/0263/001

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

Documents

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.

Labelling

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

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