FUNGICONAZOL 200 MG TABLETS FOR DOGS

Authorised

- Ketoconazole

Product identification

Medicine name:
FUNGICONAZOL 200 MG TABLETS FOR DOGS
CANIZOL VET 200 mg, tabletės šunims

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:
Lithuania

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:
Le Vet. Beheer B.V.

Marketing authorisation date:
2/11/2014

Manufacturing sites for batch release:

Genera d.d.
Lelypharma B.V.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/14/2255/001-010

Date of authorisation status change:
27/10/2019

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

13693 Final PuAR.pdf

RV2255.pdf