Finilac 50 mcg/ml

Authorised

- Cabergoline

Product identification

Medicine name:
Finilac 50 mcg/ml

Active substance:
Cabergoline

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Cabergoline
50.00 microgram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02CB03

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Croatia

Available in:
Croatia

Package description:
Available only in Croatian
Available only in Croatian
Available only in Croatian
Available only in Croatian
Available only in Croatian

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:
21/05/2015

Manufacturing sites for batch release:
Dreluso Pharmazeutika Dr. Elten und Sohn GmbH

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/20-01/173

Date of authorisation status change:
31/10/2024

Reference member state:
Netherlands

Procedure number:
NL/V/0188/001

Concerned member states:
Austria Belgium Croatia Cyprus Czechia Denmark Estonia Finland France Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg 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

Summary of Product Characteristics

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