CALIERGOLIN 50 microgram/ml oral solution for dogs and cats

Authorised

- Cabergoline

Product identification

Medicine name:
CALIERGOLIN 50 microgram/ml oral solution for dogs and cats
Caliergolin 50 microgramas/ml solução oral para cães e gatos

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

Withdrawal period by route of administration:

Oral use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02CB03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Package description:

Amber glass bottles type II (30 ml capacity) containing 24 ml of solution, closed with a grey bromobutyl rubber stopper coated with a thin fluoropolymer layer and aluminium cap, supplied with bottle adapter and PP/HDPE dosing syringe of 3 ml, packaged in Cardboard box
Amber glass bottles type III (15 ml capacity) containing 15 ml of solution, closed with a grey bromobutyl rubber stopper coated with a thin fluoropolymer layer and aluminium cap, supplied with bottle adapter and PP/HDPE dosing syringe of 3 ml, packaged in Cardboard box
Amber glass bottles type III (15 ml capacity) containing 7 ml of solution, closed with a grey bromobutyl rubber stopper coated with a thin fluoropolymer layer and aluminium cap, supplied with bottle adapter and PP/HDPE dosing syringe of 1 ml, packaged in Cardboard box

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:
Calier Portugal Medicamentos E Produtos Veterinarios S.A.

Marketing authorisation date:
27/12/2016

Manufacturing sites for batch release:
Laboratorios Calier S.A.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
1069/01/16DFVPT

Date of authorisation status change:
27/12/2016

Reference member state:
Portugal

Procedure number:
PT/V/0126/001

Concerned member states:
Italy Spain

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

Combined File of all Documents

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