Kelactin 50 µg/ml oral solution

Authorised

- Cabergoline

Product identification

Medicine name:
Kelactin 50 µg/ml oral solution

Active substance:
Cabergoline

Target species:
Cat
Dog

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:
Iceland

Package description:
Kelactin 50 µg/ml or. sol. 24 ml (glass type II)
Kelactin 50 µg/ml or. sol. 14 ml (glass type III)
Kelactin 50 µg/ml or. sol. 7 ml (glass type III)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Veyx Pharma GmbH

Marketing authorisation date:
24/02/2012

Manufacturing sites for batch release:
Veyx Pharma GmbH

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
IS/2/12/001/01

Date of authorisation status change:

16/03/2017

Reference member state:
Belgium

Procedure number:
BE/V/0025/001

Concerned member states:
Cyprus Czechia Denmark Finland France Germany Greece Hungary Iceland Italy Luxembourg Netherlands Norway Poland Portugal Slovakia Spain Sweden United Kingdom (Northern Ireland)

Generic of:
600000085401

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

Documents

Summary of Product Characteristics
Package Leaflet
Labelling