

Kelactin 50 µg/ml oral solution

Authorised

- Cabergoline

Product identification

Medicine name:

Kelactin 50 µg/ml oral solution

Kelactin Vet. 50 mikrogram/ml oral opløsning

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

Withdrawal period by route of administration:

Oral use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02CB03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

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:

12/03/2012

Manufacturing sites for batch release:

Veyx Pharma GmbH

Responsible authority:

Danish Health And Medicines Authority

Authorisation number:

48957

Date of authorisation status change:

12/03/2012

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

Package Leaflet

English (PDF)

Published on: 16/03/2023

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Summary of Product Characteristics

English (PDF)

Published on: 16/03/2023

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Labelling

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