

Kabergovet 50 microgram/ml oral solution for dogs and cats

Authorised

- Cabergoline

Product identification

Medicine name:

Kabergovet 50 microgram/ml oral solution for dogs and cats

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 liquid

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:

Ireland

Available in:

Ireland

Package description:

Cardboard box with 1 x amber PET bottle of 15 ml capacity (containing 15 ml) closed by HDPE screw cap with LDPE plug and safety seal supplied with 3 ml PP oral syringe with HDPE plunger

Cardboard box with 1 x amber PET bottle of 15 ml capacity (containing 7 ml) closed by HDPE screw cap with LDPE plug and safety seal supplied with 3 ml PP oral syringe with HDPE plunger

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:

Vet-Agro Multi-Trade Company Sp. z o.o.

Marketing authorisation date:

5/02/2021

Manufacturing sites for batch release:

Multi-Trade Company "Vet-Agro" Sp. z o.o.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA20742/007/001

Date of authorisation status change:

5/02/2021

Reference member state:

Netherlands

Procedure number:

NL/V/0312/001

Concerned member states:

Belgium Bulgaria Czechia France Greece Hungary Ireland Italy Lithuania
Poland Portugal Spain

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

Source URL: <https://medicines.health.europa.eu/veterinary/600000032039>