File downloaded on 2026-11-26
Source URL: https://medicines.health.europa.eu/veterinary/en/600000032039

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

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:
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Marketing authorisation date:
5/02/2021

Manufacturing sites for batch release:
Przedsiebiorstwo Wielobranzowe 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