Pergoquin, 1 mg tablets for horses

Authorised

- Pergolide

Product identification

Medicine name:
Pergoquin, 1 mg tablets for horses
Pergoquin 1 mg tablety pre kone

Active substance:
Pergolide

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Pergolide
1.00 milligram(s) / 1.00 Piece

Pharmaceutical form:
Tablet

Withdrawal period by route of administration:

Oral use:

- **Horse**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN04BC02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
Cardboard box of 200 tablets containing Aluminium-OPA, Aluminium, PVC blisters containing 10 tablets.
Cardboard box of 160 tablets containing Aluminium-OPA, Aluminium, PVC blisters containing 10 tablets.
Cardboard box of 150 tablets containing Aluminium-OPA, Aluminium, PVC blisters containing 10 tablets.
Cardboard box of 100 tablets containing Aluminium-OPA, Aluminium, PVC blisters containing 10 tablets.
Cardboard box of 50 tablets containing Aluminium-OPA, Aluminium, PVC blisters containing 10 tablets.
Cardboard box of 60 tablets containing Aluminium-OPA, Aluminium, PVC blisters containing 10 tablets.

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:

Vetviva Richter GmbH

Marketing authorisation date:
16/12/2019

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/052/DC/19-S

Date of authorisation status change:
16/12/2019

Reference member state:
Netherlands

Procedure number:
NL/V/0295/001

Concerned member states:
Austria Belgium Bulgaria Croatia Czechia Denmark Estonia Finland Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden
United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)

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