Prasequine 1 mg tablets for horses

Authorised

- Pergolide

Product identification

Medicine name:
Prasequine 1 mg tablets for horses
Prasequin vet. 1 mg tabletki

Active substance:
Pergolide

Target species:
Horse (non food-producing)

Route of administration:
Oral use

Product details

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

Pharmaceutical form:
Tablet

Withdrawal period by route of administration:

Oral use:

- **Horse (non food-producing)**
  - Not applicable. no withdrawal period

Not authorised for use in horses intended for human consumption. The horse must have been declared as not intended for human consumption under national horse passport legislation. Not authorised for use in mares producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN04BC02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Poland

Package description:

Carton box containing 13 OPA/aluminium/PVC-aluminium blisters, containing 7 tablets each

Carton box containing 24 OPA/aluminium/PVC-aluminium blisters, containing 10 tablets each

Carton box containing 16 OPA/aluminium/PVC-aluminium blisters, containing 10 tablets each

Carton box containing 10 OPA/aluminium/PVC-aluminium blisters, containing 10 tablets each

Carton box containing 6 OPA/aluminium/PVC-aluminium blisters, containing 10 tablets each

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Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic (abridged application) - art 13(1)

Marketing authorisation holder:
CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:
27/01/2023

Manufacturing sites for batch release:
CP-Pharma Handelsgesellschaft mbH

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3239

Date of authorisation status change:
27/01/2023

Reference member state:
Netherlands

Procedure number:
NL/V/0368/001

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

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.