

# Prascend 1 mg tablets for horses

Authorised

- Pergolide mesilate

## Product identification

**Medicine name:**

Prascend 1 mg tablets for horses

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**Active substance:**

Pergolide mesilate

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**Target species:**

Horse (non food-producing)

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Pergolide mesilate

1.31 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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**Withdrawal period by route of administration:**

**Oral use:**

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**Horse (non food-producing)**

- Meat and offal. 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.

- Milk. 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.

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QN04BC02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Norway

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**Available in:**

Norway

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**Package description:**

(ID1) 100 Tablet: Box (cardboard) with 1 Blister (aluminium; polyvinyl chloride; nylon) with 100 Tablet

(ID2) 60 Tablet: Box (cardboard) with 1 Blister (aluminium; polyvinyl chloride; nylon) with 60 Tablet

(ID3) 160 Tablet: Box (cardboard) with 1 Blister (aluminium; polyvinyl chloride; nylon) with 160 Tablet

(ID4) 91 Tablet: Box (cardboard) with 13 Blister (nylon; aluminium; polyvinyl chloride) each with 7 Tablet, closed with Foil (vinyl, aluminium)

(ID5) 480 Tablet: Box (cardboard) with 3 Box (cardboard) each with 16 Blister (nylon; aluminium; polyvinyl chloride) each with 10 Tablet, closed with Foil (vinyl, aluminium)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Boehringer Ingelheim Vetmedica GmbH

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**Marketing authorisation date:**

20/06/2012

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Vetmedica GmbH  
Haupt Pharma Amareg GmbH

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**Responsible authority:**

Norwegian Medical Products Agency

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**Authorisation number:**

11-8625

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**Date of authorisation status change:**

29/07/2014

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0130/001

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**Concerned member states:**

Austria Belgium Denmark Finland France Iceland Ireland Italy Luxembourg  
Netherlands Norway Sweden United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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.

Combined File of all Documents

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