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Prascend 1 mg tablets for horses

- Pergolide mesilate

Authorised

# Product identification

Medicine name:
Prascend 1 mg tablets for horses
Active substance:

- Pergolide mesilate

Target species:

- Horse (non-food-producing)

Route of administration:

- Oral use

# Product details

Active substance and strength:

- Pergolide mesilate
  1.31
  milligram(s)
  /
  1.00
  Tablet

Pharmaceutical form:

- Tablet

Withdrawal period by route of administration:

- Oral use
  - 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.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

- QN04BC02

Legal status of supply:

- Veterinary medicinal product subject to veterinary prescription

Authorisation status:

- Valid

Authorised in:

- Germany

Available in:

- Germany

Package description:

- (ID5) 480 Tablet: Box (Cardboard) with 3 Box (Cardboard) each with 16 Blister (PolyAmide; Aluminium; PolyVinyl Chloride) each with 10 Tablet, closed with Foil (vinyl, Aluminium)
- (ID4) 91 Tablet: Box (Cardboard) with 13 Blister (PolyAmide; Aluminium; PolyVinyl Chloride) each with 7 Tablet, closed with Foil (vinyl, Aluminium)
- (ID3) 160 Tablet: Box (Cardboard) with 1 Blister (Aluminium; PolyVinyl Chloride; PolyAmide) with 160 Tablet
- (ID2) 60 Tablet: Box (Cardboard) with 1 Blister (Aluminium; PolyVinyl Chloride; PolyAmide) with 60 Tablet
- (ID1) 100 Tablet: Box (Cardboard) with 1 Blister (Aluminium; PolyVinyl Chloride; PolyAmide) with 100 Tablet

# Additional information

Entitlement type:

- Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

- Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:

- 20/08/2009

Manufacturing sites for batch release:

- Boehringer Ingelheim Vetmedica GmbH
- Haupt Pharma Amareg GmbH

Responsible authority:

- Federal Office Of Consumer Protection And Food Safety

Authorisation number:

- 401219.00.00

Date of authorisation status change:

- 29/10/2014

Reference member state:

- Germany

Procedure number:

- DE/V/0130/001

Concerned member states:

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

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

# Documents

# Product information

Combined File of all Documents

English (PDF)
Published on: 27/03/2026
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Other languages (1)
German (PDF)
Published on: 14/03/2026
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# Public assessment report(s)

2401219-paren-20200625.pdf

English (PDF)
Published on: 27/03/2026
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