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# Prid delta 1.55 g vaginal delivery system for cattle

Authorised

- Progesterone

## Product identification

**Medicine name:**

Prid delta 1.55 g vaginal delivery system for cattle

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**Active substance:**

Progesterone

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**Target species:**

Cattle

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**Route of administration:**

Vaginal use

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## Product details

**Active substance and strength:**

Progesterone

1.55 gram(s) / 1.00 System

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**Pharmaceutical form:**

Vaginal delivery system

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**Withdrawal period by route of administration:****Vaginal use:**

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**Cattle**

- Meat and offal. 0 day
- Milk. 0 day

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

QG03DA04

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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:**

Ireland

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**Package description:**

Cardboard box containing 10 sachets of 1 device

Cardboard box containing 100 sachets of 1 device

Sachet containing 10 devices

Polyethylene box containing 1 applicator and 50 sachets of 1 device

Polyethylene box containing 50 sachets of 1 device

Cardboard box containing 1 applicator and 50 sachets of 1 device

Cardboard box containing 50 sachets of 1 device

Cardboard box containing 1 applicator and 25 sachets of 1 device

Cardboard box containing 25 sachets of 1 device

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ceva Sante Animale

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**Marketing authorisation date:**

10/12/2010

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**Manufacturing sites for batch release:**

Ceva Sante Animale

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10815/013/001

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**Date of authorisation status change:**

10/12/2010

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**Reference member state:**

France

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**Procedure number:**

FR/V/0215/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania  
Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain  
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

Combined File of all Documents

eu-puar-frv0215001-mr-rpe498-en.pdf