

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

Active substance:

Progesterone

Target species:

Cattle

Route of administration:

Vaginal use

Product details

Active substance and strength:

Progesterone

1.55 gram(s) / 1.00 System

Pharmaceutical form:

Vaginal delivery system

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

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Cattle

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG03DA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Available in:

Belgium

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

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:

Ceva Sante Animale

Marketing authorisation date:

26/08/2010

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V376415

Date of authorisation status change:

26/08/2010

Reference member state:

France

Procedure number:

FR/V/0215/001

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)

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.

Combined File of all Documents

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