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:

- **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:

Spain

Available in:

Spain

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 Salud Animal S.A.

Marketing authorisation date:
30/09/2010

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
2194 ESP

Date of authorisation status change:
30/09/2010

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