# CIDR 1.38 g Vaginal Delivery System for Cattle

Authorised

- Progesterone

## Product identification

**Medicine name:**
CIDR 1.38 g Vaginal Delivery System for Cattle
CIDR 1.38 g Hulpmiddel voor vaginaal gebruik
CIDR 1.38 g Système de diffusion vaginal
CIDR 1.38 g vaginales Wirkstofffreisetzungssystem

**Active substance:**
Progesterone

**Target species:**
Cattle (cow)
Cattle (heifer)

**Route of administration:**
Vaginal use

## Product details

**Active substance and strength:**
Progesterone
1.38 gram(s) / 1.00 System

**Pharmaceutical form:**
Vaginal delivery system

**Withdrawal period by route of administration:**

**Vaginal use:**

- **Cattle (cow)**
  - Meat and offal. no withdrawal period — withdrawal period is 0 days
  - Milk. no withdrawal period — withdrawal period is 0 hours
- **Cattle (heifer)**
  - Meat and offal. no withdrawal period — withdrawal period is 0 days
  - Milk. no withdrawal period — withdrawal period is 0 hours

**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:**
bag containing 10 vaginal delivery systems of 1,38 g

## 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:**
Zoetis Belgium

**Marketing authorisation date:**
10/12/2007

**Manufacturing sites for batch release:**
Zoetis Belgium

**Responsible authority:**
Federal Agency For Medicines And Health Products

**Authorisation number:**
BE-V309032

**Date of authorisation status change:**
10/12/2007

**Reference member state:**
Spain

**Procedure number:**
ES/V/0318/001

**Concerned member states:**
Austria Belgium Czechia Denmark Finland France Germany Hungary Ireland Italy Luxembourg Netherlands Norway Poland Portugal Slovakia Slovenia Sweden United Kingdom (Northern Ireland)

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

## Documents

Summary of Product Characteristics

English (PDF)
Published on: 23/03/2023
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Package Leaflet

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Labelling

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Combined File of all Documents