CIDR 1.38 g Vaginal Delivery System for Cattle

Authorised

• Progesterone

Product identification

Medicine name:

CIDR 1.38 g Vaginal Delivery System for Cattle Relmont vet 1,38 g vaginalinnlegg for storfe

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:

Norway

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 Animal Health ApS

Marketing authorisation date: 7/05/2012
Manufacturing sites for batch release: Zoetis Belgium
Responsible authority: Norwegian Medical Products Agency
Authorisation number: 10-8040
Date of authorisation status change: 24/10/2012
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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