# RILEXINE DC 375 MG INTRAMAMMARY SUSPENSION FOR DRY COWS

Authorised

- Cephalexin benzathine

## Product identification

**Medicine name:**
RILEXINE DC 375 MG INTRAMAMMARY SUSPENSION FOR DRY COWS
Rilexine DC, 375 mg, intramaminė suspensija užtrūkusioms karvėms

**Active substance:**
Cephalexin benzathine

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

**Route of administration:**
Intramammary use

## Product details

**Active substance and strength:**
Cephalexin benzathine
500.00 milligram(s) / 1.00 Syringe

**Pharmaceutical form:**
Intramammary suspension

**Withdrawal period by route of administration:**

**Intramammary use:**

- **Cattle (dry cow)**
  - Meat and offal. 4 day — Meat and offal: 4 days
  - Milk. 43 day — 42.5 days after treatment when dry period is 42 days or less
  - Milk. 12 hour — 12 hours after calving when dry period is more than 42 days

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QJ51DB01

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Lithuania

**Available in:**
Lithuania

**Package description:**
Box of 12 x 8g intramammary syringes and 12 cleaning towels
Box of 60 x 8g intramammary syringes and 60 cleaning towels
Box of 24 x 8g intramammary syringes and 24 cleaning towels

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Virbac

**Marketing authorisation date:**
5/04/2022

**Manufacturing sites for batch release:**
Haupt Pharma Latina S.r.l.
Virbac

**Responsible authority:**
State Food And Veterinary Service

**Authorisation number:**
LT/2/22/2703/001-003

**Date of authorisation status change:**
5/04/2022

**Reference member state:**
France

**Procedure number:**
FR/V/0438/001

**Concerned member states:**
Austria Belgium Bulgaria Croatia Czechia Denmark Estonia Finland Germany Hungary Ireland Italy Latvia Lithuania 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

RV2703.pdf