

# Arentor DC 250 mg Intramammary Suspension for Dry Cows

Authorised

- Cefalonium dihydrate

## Product identification

**Medicine name:**

Arentor DC 250 mg Intramammary Suspension for Dry Cows

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**Active substance:**

Cefalonium dihydrate

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**Target species:**

Cattle

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**Route of administration:**

Intramammary use

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## Product details

**Active substance and strength:**

Cefalonium dihydrate

269.63 milligram(s) / 1.00 Syringe

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**Pharmaceutical form:**

Intramammary suspension

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**Withdrawal period by route of administration:**

**Intramammary use:**

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**Cattle**

- Meat and offal. 21 day

- Milk. 96 hour 96 hours after calving if the dry period is longer than 54 days

- Milk. 58 day

58 days following the treatment if the dry period is less than or equal to 54 days

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

QJ51DB90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Ireland

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**Package description:**

A 3g coloured low density polyethylene intramammary syringe with a coloured low density polyethylene dual capPack sizes: Buckets of 120 intramammary syringes and 120 individually wrapped cleaning towels containing isopropyl alcohol.

A 3g coloured low density polyethylene intramammary syringe with a coloured low density polyethylene dual capPack sizes: Cartons of 20 intramammary syringes and 20 individually wrapped cleaning towels containing isopropyl alcohol.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Univet Limited

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**Marketing authorisation date:**

7/12/2018

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**Manufacturing sites for batch release:**

Univet Limited

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10990/050/001

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**Date of authorisation status change:**

7/12/2018

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0389/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia France Germany  
Greece Hungary Italy Latvia Lithuania Luxembourg Netherlands Poland  
Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics